



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

<p>(51) International Patent Classification ⁶ : C07C 233/65, A01N 37/22, C07C 235/42, 235/84, 63/68</p>	A1	<p>(11) International Publication Number: WO 98/45252</p> <p>(43) International Publication Date: 15 October 1998 (15.10.98)</p>		
<table style="width: 100%; border: none;"> <tr> <td style="width: 50%; vertical-align: top; padding: 5px;"> <p>(21) International Application Number: PCT/EP98/02014</p> <p>(22) International Filing Date: 7 April 1998 (07.04.98)</p> <p>(30) Priority Data: 97/04322 9 April 1997 (09.04.97) FR</p> <p>(71) Applicant (for all designated States except US): HOECHST SCHERING AGREVO S.A. [FR/FR]; 163, avenue Gambetta, F-75020 Paris (FR).</p> <p>(72) Inventors; and (75) Inventors/Applicants (for US only): BROUILLARD, Agnès [FR/FR]; 14, rue Gounod, F-93250 Villemomble (FR). DEMASSEY, Jacques [FR/FR]; 29, allée Saint Jacques Chalifert, F-77144 Montevrain (FR). DUTHEIL, Philippe [FR/FR]; 96, rue Guy Mocquet, F-94340 Joinville le Pont (FR). WESTON, John [GB/FR]; 7, résidence Desaix Av. Desaix Pav. 24, F-78600 Maison Lafitte (FR).</p> <p>(74) Agent: RIPPEL, Hans, Christoph; Hoechst Schering AgrEvo GmbH, Patent- und Lizenzabteilung, Gebäude K 801, D-65926 Frankfurt am Main (DE).</p> </td> <td style="width: 50%; vertical-align: top; padding: 5px;"> <p>(81) Designated States: AL, AM, AU, AZ, BA, BB, BG, BR, BY, CA, CN, CU, CZ, EE, GE, GW, HU, ID, IL, IS, JP, KG, KP, KR, KZ, LC, LK, LR, LT, LV, MD, MG, MK, MN, MX, NO, NZ, PL, RO, RU, SG, SI, SK, SL, TJ, TM, TR, TT, UA, US, UZ, VN, YU, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).</p> <p>Published <i>With international search report.</i></p> </td> </tr> </table>			<p>(21) International Application Number: PCT/EP98/02014</p> <p>(22) International Filing Date: 7 April 1998 (07.04.98)</p> <p>(30) Priority Data: 97/04322 9 April 1997 (09.04.97) FR</p> <p>(71) Applicant (for all designated States except US): HOECHST SCHERING AGREVO S.A. [FR/FR]; 163, avenue Gambetta, F-75020 Paris (FR).</p> <p>(72) Inventors; and (75) Inventors/Applicants (for US only): BROUILLARD, Agnès [FR/FR]; 14, rue Gounod, F-93250 Villemomble (FR). DEMASSEY, Jacques [FR/FR]; 29, allée Saint Jacques Chalifert, F-77144 Montevrain (FR). DUTHEIL, Philippe [FR/FR]; 96, rue Guy Mocquet, F-94340 Joinville le Pont (FR). WESTON, John [GB/FR]; 7, résidence Desaix Av. Desaix Pav. 24, F-78600 Maison Lafitte (FR).</p> <p>(74) Agent: RIPPEL, Hans, Christoph; Hoechst Schering AgrEvo GmbH, Patent- und Lizenzabteilung, Gebäude K 801, D-65926 Frankfurt am Main (DE).</p>	<p>(81) Designated States: AL, AM, AU, AZ, BA, BB, BG, BR, BY, CA, CN, CU, CZ, EE, GE, GW, HU, ID, IL, IS, JP, KG, KP, KR, KZ, LC, LK, LR, LT, LV, MD, MG, MK, MN, MX, NO, NZ, PL, RO, RU, SG, SI, SK, SL, TJ, TM, TR, TT, UA, US, UZ, VN, YU, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).</p> <p>Published <i>With international search report.</i></p>
<p>(21) International Application Number: PCT/EP98/02014</p> <p>(22) International Filing Date: 7 April 1998 (07.04.98)</p> <p>(30) Priority Data: 97/04322 9 April 1997 (09.04.97) FR</p> <p>(71) Applicant (for all designated States except US): HOECHST SCHERING AGREVO S.A. [FR/FR]; 163, avenue Gambetta, F-75020 Paris (FR).</p> <p>(72) Inventors; and (75) Inventors/Applicants (for US only): BROUILLARD, Agnès [FR/FR]; 14, rue Gounod, F-93250 Villemomble (FR). DEMASSEY, Jacques [FR/FR]; 29, allée Saint Jacques Chalifert, F-77144 Montevrain (FR). DUTHEIL, Philippe [FR/FR]; 96, rue Guy Mocquet, F-94340 Joinville le Pont (FR). WESTON, John [GB/FR]; 7, résidence Desaix Av. Desaix Pav. 24, F-78600 Maison Lafitte (FR).</p> <p>(74) Agent: RIPPEL, Hans, Christoph; Hoechst Schering AgrEvo GmbH, Patent- und Lizenzabteilung, Gebäude K 801, D-65926 Frankfurt am Main (DE).</p>	<p>(81) Designated States: AL, AM, AU, AZ, BA, BB, BG, BR, BY, CA, CN, CU, CZ, EE, GE, GW, HU, ID, IL, IS, JP, KG, KP, KR, KZ, LC, LK, LR, LT, LV, MD, MG, MK, MN, MX, NO, NZ, PL, RO, RU, SG, SI, SK, SL, TJ, TM, TR, TT, UA, US, UZ, VN, YU, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).</p> <p>Published <i>With international search report.</i></p>			
<p>(54) Title: AROMATIC AMIDES, THEIR PREPARATION PROCESS AND THEIR USE AS PESTICIDES</p> <p>(57) Abstract</p> <p style="margin-left: 20px;">Aromatic amides, their preparation process and their use as pesticides.</p> <div style="text-align: center; margin-top: 20px;"> </div>				

FOR THE PURPOSES OF INFORMATION ONLY

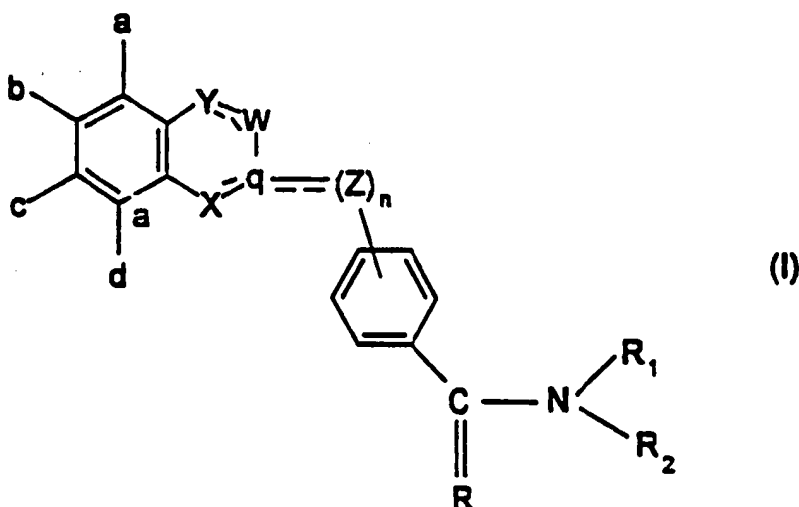
Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece			TR	Turkey
BG	Bulgaria	HU	Hungary	ML	Mali	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MN	Mongolia	UA	Ukraine
BR	Brazil	IL	Israel	MR	Mauritania	UG	Uganda
BY	Belarus	IS	Iceland	MW	Malawi	US	United States of America
CA	Canada	IT	Italy	MX	Mexico	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NE	Niger	VN	Viet Nam
CG	Congo	KE	Kenya	NL	Netherlands	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NO	Norway	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	NZ	New Zealand		
CM	Cameroon			PL	Poland		
CN	China	KR	Republic of Korea	PT	Portugal		
CU	Cuba	KZ	Kazakhstan	RO	Romania		
CZ	Czech Republic	LC	Saint Lucia	RU	Russian Federation		
DE	Germany	LI	Liechtenstein	SD	Sudan		
DK	Denmark	LK	Sri Lanka	SE	Sweden		
EE	Estonia	LR	Liberia	SG	Singapore		

Aromatic amides, their preparation process and their use as pesticides

The present invention relates to aromatic amides, their preparation process and their use as pesticides.

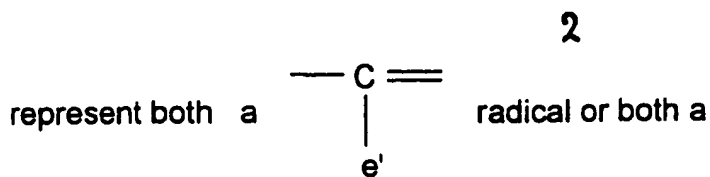
A subject of the invention is the compounds of formula (I):



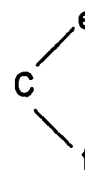
in which:

- a, b, c and d, identical to or different from one another, represent a hydrogen atom, a halogen atom, an alkyl, alkenyl or alkynyl, O-alkyl, O-alkenyl or O-alkynyl, S-alkyl, S-alkenyl or S-alkynyl radical containing up to 8 carbon atoms, optionally substituted by one or more halogen atoms, a C≡N, NO₂ or NH₂ radical, the substituents a, b, c and d being able to form between themselves rings, preferably one, which either contain or do not contain one or more, preferably one or two, hetero atoms, preferably from the group consisting of S, O and N, and which are substituted, or unsubstituted.

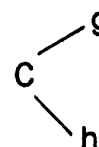
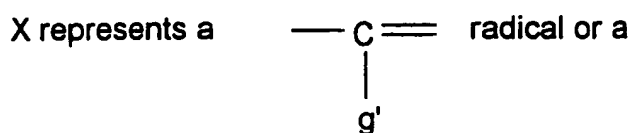
- Y and W, identical to or different from one another,



2



radical in which e, f and e', identical or different, represent a hydrogen atom, a halogen atom, a free, etherified or esterified hydroxyl radical, or an alkyl radical containing up to 8 carbon atoms optionally substituted by one or more halogen atoms;

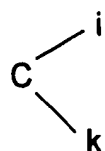


radical in which g, h and g', identical or different, represent a hydrogen atom, a halogen atom, a free, etherified or esterified hydroxyl radical, or an alkyl radical containing up to 8 carbon atoms optionally substituted by one or more halogen atoms, or X represents a C=O radical, an oxygen atom or a nitrogen atom, or X forms with the carbon in position 2 belonging to radical q an epoxy bridge, a cyclic hydrocarbonated radical optionally substituted by one or more halogen atoms;

- q represents a C= radical or a CD radical, in which D represents a hydrogen atom, a halogen atom or an alkyl or alkoxy radical containing up to 8 carbon atoms optionally substituted by one or more halogen atoms, or D forms with the carbon atom which carries it and one of the carbon atoms adjacent to it a carbon-carbon double bond, an epoxy radical, a cyclic hydrocarbonated radical, optionally substituted by one or more halogen atoms;

n represents an integer varying from 0 to 8;

- Z represents a



radical in which i and k, identical

3

or different, represent a hydrogen atom, a halogen atom, a $C\equiv N$ radical, a free, etherified or esterified hydroxyl radical, an SR' radical, wherein R' is an organic rest containing up to 8 carbon atoms, preferably S-alkyl, or an alkyl radical containing up to 8 carbon atoms optionally substituted by one or more halogen atoms, or Z represents an oxygen, sulfur, nitrogen atom, a $C=O$ or $C=S$ radical, it being understood that if n is greater than 1, Z can take different values,

- R represents an oxygen or sulfur atom;
 - R_1 and R_2 , identical to or different from one another, represent a hydrogen atom, a linear, branched or cyclic, saturated or unsaturated, optionally substituted alkyl, CO-alkyl, CONH-alkyl or CO_2 alkyl radical containing up to 8 carbon atoms, optionally interrupted by one or more preferably non-adjacent heteroatoms, preferably from the group consisting of N, O, S, or an optionally substituted aryl or heteroaryl radical,
 - the $-C-(Z)_n$ chain is fixed in position 3 or 4 of the benzamide, the dotted lines representing one or more optional double bonds,
- in all their possible isomeric forms as well as their mixtures.

By compound of formula (I) are designated all the geometric isomers and stereoisomers taken individually or in a mixture.

In the definition of substituents:

- alkyl preferably represents a methyl, ethyl, propyl, isopropyl, butyl, isobutyl,
- n-pentyl, isopentyl, cyclopropyl, cyclobutyl or cyclopentyl radical,
- alkenyl preferably represents a vinyl, 1-propenyl, 2-methyl 2-propenyl or isopropenyl radical,
- alkynyl preferably represents an ethynyl, 1-propynyl, 2-propynyl or pent-2-ene-4-enyl radical,
- halogen preferably represents a fluorine, chlorine, bromine or iodine atom,
- aryl preferably represents a carbocyclic aromatic group containing 4 to 10 carbon atoms, particularly preferably a phenyl, naphthyl, dihydronaphthyl, tetrahydronaphthyl, indanyl or indenyl radical,

4

- a heterocyclic radical is preferably a heteroaryl radical or a saturated or unsaturated 3 to 8 membered ring comprising one, two three or four heteroatoms from the group consisting of N, O and S.
- heteroaryl is preferably a 3 to 7 membered aromatic ring comprising one, two, three or four heteroatoms from the group consisting of N, O and S, particularly preferred thienyl, furyl, pyrrolyl, imidazolyl, pyrazolyl, pyridyl, pyrazinyl, pyrimidinyl, pyridazinyl, isothiazolyl, isoxazolyl, oxazolyl, thiazolyl, oxadiazolyl and tetrazinyl
- Particularly preferred heterocyclic radicals are a thienyl, furyl, pyrannyl, pyrrolyl, 2H-pyrrolyl, imidazolyl, pyrazolyl, pyridyl, pyrazinyl, pyrimidinyl, pyridazinyl, isothiazolyl, isoxazolyl, furazanyl, thiazolyl, oxazolyl, pyrrolidinyl, pyrrolinyl, imidazolidinyl, pyrazolinyl, piperidyl, piperazinyl, morpholinyl, izapinyl, thiazinyl, tetrazinyl, oxathiolanyl or thiadiazinyl radical.

When the aryl or heterocyclic radical is substituted, it is preferably substituted by one or more substitutents chosen in particular from halogen atoms, alkyl or alkoxy radicals containing up to 8 carbon atoms, or methylenedioxy, difluoromethylenedioxy, tetrafluoro ethylenedioxy, cyano, nitro, cyanato, thiocyanato, pentafluorothio or fluorosulfonyl groups.

Etherified or esterified preferably means etherified with a linear or branched C₁-C₈-alkyl group or esterified with a (C₁-C₈)-carboxylic acid.

- If any of the substituents a-d form a ring it is preferably a 4 to 8 membered ring which is preferably monounsaturated (due to fusion with the naphthyl group) and is carbocyclic or contains preferably one or two heteroatoms from the group consisting of N, O and S.
- If Z is O, N, or CO, CS and n is greater then 1 it is preferred that heteroatoms are not adjacent to each other and CO, CS are not adjacent to each other.
- It goes without saying that x, q and Z have to be chosen in a way that a tetravalent carbon at q results.

5

Particularly preferred are compounds of formula (I) in which Y represents a $\text{-CH}_2\text{-}$ radical, those in which Y and W represent a CH radical and together form a double bond in position 3(4), those in which W represents a CH_2 radical, those in which q and X represent a CH= radical and together form a double bond, those in which q represents a CH or C= radical, those in which X represents a CH, a CH_2 , a CHOH or a CO radical, those in which Z represents a CH_2 , a CHF, a CHOH or a COOCH_3 radical, those in which n represents the number 1, those in which R represents an oxygen atom, those in which R_1 represents a hydrogen atom, those in which R_2 represents an alkyl radical containing up to 8 carbon atoms or a phenyl radical optionally substituted by one or more halogen atoms and/or one or more linear or branched alkyl radicals containing up to 8 carbon atoms in particular in which R_2 represents an alkyl radical containing up to 6 carbon atoms and in particular an isobutyl radical or a 2-methylphenyl radical, those in which at least one of substituents a, b, c and d represents a halogen atom, $(\text{C}_1\text{-C}_8)$ alkyl or $(\text{C}_1\text{-C}_8)$ alkoxy for example those in which two of the substituents a, b, c and d represent a chlorine or bromine atom, the compounds of formula (I) in which two of the substituents a, b, c and d represent a hydrogen atom.

In particular a subject of the invention is the compounds the preparation of which is given hereafter in the experimental part and quite particularly the compounds of Examples A, B, C, D, E, F, G, H, I, J, K, L and M.

The compounds of formula (I) can be used to combat harmful organisms such as arthropods, for example insects and acaridae, and helminths, for example nematodes, or molluscs, for example slugs. Therefore a subject of the present invention is a process for combating arthropods and/or helminths and/or molluscs, which comprises the administration to the arthropods and/or helminths and/or molluscs, or to their environment, of a quantity of a compound of formula (I) which is sufficient to destroy the harmful organism. Also a subject of the present invention is a process for combating and/or eradicating infestations by arthropods and/or helminths and/or molluscs of animals (including humans) and/or of plants (including trees) and/or stored products, which comprises the administration to the animal or to the locality of an effective quantity of a compound of formula (I). A subject of the

6

invention is also the compounds of formula (I) to be used in human and veterinary medicine, in public health and/or in agriculture for combating harmful arthropods and/or helminths.

The compounds of formula (I) are particularly valuable in the protection of standing crops, forage crops, crops in plantations, in greenhouses, in orchards and in vineyards, of ornamental plants and trees in plantations and forests, for example cereals (such as corn, wheat, rice, sorghum), cotton, tobacco, vegetables and salad vegetables (such as beans, cabbages, cucurbitaceae, lettuces, onions, tomatoes and peppers), food crops (such as potatoes, sugar beet, peanuts, soya, oilseed rape), sugar cane, meadows and forage (such as corn, sorghum, alfalfa), plantations (such as those producing tea, coffee, cocoa, banana, palm oil, coconut, rubber, spices), orchards and tree plantations (such as those producing stone fruits and pome fruit, citrus fruits, kiwis, avocados, mangoes, olives and walnuts), vines, ornamental plants, flowers and bushes in greenhouses and in gardens and parks, forest trees (both deciduous and evergreen) in forests, plantations and nurseries. They are also valuable in the protection of timber (standing, felled, processed, stored or in buildings) against attack from wood wasps (for example *Urocerus*) or coleopterous insects (for example *scolytidae*, *platypodidae*, *lyctidae*, *bostrichidae*, *cerambycidae*, *anobiidae*) and termites.

They can be used in the protection of stored products, such as grains, fruits, nuts, spices and tobacco, whether whole, ground or converted into products, against attack from mites, coleopterous insects and weevils. Stored animal products such as skins, furs, wool and feathers, in natural or processed form (for example rugs or textile materials) can also be protected against attack from mites and coleopterous insects; similarly meat and fish can be protected against attack from coleopterous insects and flies.

The compounds of general formula (I) are particularly useful for combating arthropods, helminths or molluscs, which are harmful to man and domestic animals, or spread or are carriers of diseases affecting the latter, for example those which

have been described above, and more particularly for combating ticks, mites, lice, fleas, midges and flies which cause bites and are harmful.

The invention also relates to the use of the compounds of formula (I) as defined previously, as pesticides in particular as insecticides, acaricides and nematocides in the protection of crops in particular rice and cotton crops, or for the treatment of premises for storing said crops and in particular as insecticides and acaricides in domestic or public premises.

The compounds of formula (I) can be used to these ends by the application of the compounds as they are, or in a diluted form, in a known manner, in the form of dips, sprays, mists, lacquers, foams, powders, dusting products, aqueous suspensions, pastes, gels, shampoos, ointments, combustible solids, spray pads, combustible coils, baits, food additives, wettable powders, granules, aerosols, emulsifiable concentrates, oily suspensions, oily solutions, pressurized sprays, impregnated articles, lotions or other standard compositions well known to a person skilled in the art. Concentrates for dips are not used as they are, but diluted with water, and the animals are immersed in a dipping bath containing the dipping product. Sprays can be applied by hand, or by means of a spray lance or frame. The animal, the ground, the plant or the surface can be saturated with the spray using a high volume application, or coated superficially by spraying with a light or very low-volume application. Aqueous suspensions can be applied to the animal in the same manner as sprays or dips. Dusting products must be distributed via a powder applicator or, in the case of animals, be incorporated in perforated bags fixed to trees or poles. Pastes, shampoos and ointments can be applied by hand or spread on the surface of an inert material against which the animals rub themselves and thus transfer the product onto their skin. Lotions are distributed as low-volume amounts of liquid on the backs of animals, so that all or most of the liquid remains on the animals.

The compounds of formula (I) can be presented as ready-to-use compositions for use on plants, animals or surfaces, or in the form of compositions which need to be diluted before use, but both types of compositions contain a compound of formula (I) intimately mixed with one or more excipients or diluents. The excipients can be

8

liquid, solid or gaseous, or can contain mixtures of such substances, and the compound of formula (I) can be present in a concentration of 99 to 0.025 % w/v., according to whether the composition needs a stronger or weaker dilution.

Dusting products, powders and granules contain the compound of formula (I) intimately mixed with a pulverulent solid inert excipient, for example suitable clays, kaolin, bentonite, attapulgit, adsorbent carbon black, talc, mica, chalk, gypsum, tricalcium phosphate, powdered cork, magnesium silicate, vegetable excipients, starch and diatomaceous earths. These solid compositions are in general prepared by impregnating the solid diluents with solutions of the compound of formula (I) in volatile solvents, by evaporating the solvents and, if appropriate, grinding of the products to obtain powders and, if desired, by granulating, compacting or encapsulating the products.

The sprays of a compound of formula (I) can contain a solution in an organic solvent (for example those mentioned above) or an emulsion in water (dipping or spraying), prepared on site from an emulsifiable concentrate (otherwise known as oil miscible with water), which can also be used for dipping. The concentrate preferably contains a mixture of the active ingredient, with or without organic solvent, and one or more emulsifiers. Solvents can be present in very variable quantities, but preferably in a quantity of 0 to 90% w/v of the composition, and can be chosen from kerosene, ketones, alcohols, xylene, aromatic naphtha, and other solvents known for use in compositions. The concentration of emulsifiers can be very variable, but is preferably in the range of 5 to 25% w/v, and the emulsifiers are advantageously non-ionic surfactants, in particular polyoxy-alkylenic esters of alkylphenols and polyoxyethylenic derivatives of hexitol anhydrides or anionic surfactants, in particular sodium laurylsulfate, fatty alcohol ethersulfates, the sodium and calcium salts of alkylarylsulfonates and alkylsulfo-succinates.

The cationic emulsifiers are in particular benzalkonium chloride and quaternary ammonium ethylsulfates.

The amphoteric emulsifiers are in particular carboxymethylated oleic imidazoline and alkyldimethyl-betaines.

Vaporization wicks normally contain a mixture of cotton and cellulose compressed into a pad, e.g. of approximately 32 mm by 22 mm by 3 mm, treated with, normally, up to 0.3 ml of a concentrate which contains the active ingredient in an organic solvent and optionally an anti-oxidant, a coloring agent and a perfume.

The insecticide may be vaporized by using a heat source, such as an electrically-powered device for heating the wicks.

The combustible solids normally contain sawdust and a binder mixed with the active ingredient and used in the form of molded strips (usually in coils). A coloring agent and a fungicide can also be added.

The wettable powders contain an inert solid excipient, one or more surfactants, and optionally stabilizers and/or anti-oxidants.

The emulsifiable concentrates contain emulsifying agents and often an organic solvent, such as kerosene, ketones, alcohols, xylenes, aromatic naphtha and other known solvents.

The wettable powders and emulsifiable concentrates normally contain 5 to 95% by weight of the active ingredient and are diluted, for example with water, before use.

The lacquers contain a solution of the active ingredient in an organic solvent, together with a resin and optionally a plastizer.

Dips can be prepared not only from emulsifiable concentrates, but also from wettable powders, dips based on soap and aqueous suspensions containing a compound of formula (I) intimately mixed with a dispersing agent and one or more surfactants.

The aqueous suspensions of a compound of formula (I) can include a suspension in water together with suspension agents, stabilizers or other agents. The

10

suspensions or solutions can be applied as they are or in a form diluted in a known manner.

The ointments (or greases) can be prepared from vegetable oils, synthetic esters of fatty acids or lanolin, together with an inert base such as soft paraffin. A compound of formula (I) is preferably distributed uniformly throughout the mixture, in solution or in suspension. Ointments can also be obtained from emulsifiable concentrates by dilution of the latter with an ointment base.

The pastes and shampoos are also semi-solid compositions in which a compound of formula (I) can be present in uniform dispersion in a suitable base such as soft or liquid paraffin, or in a non-fat base with glycerol, a glue or a suitable soap. Since the ointments, shampoos and pastes are normally applied without any other dilution, they must contain the appropriate percentage of the compound of formula (I) required for the treatment.

Aerosol sprays can be prepared in the form of a simple solution of the active ingredient in the aerosol propellant and the co-solvent, such as a halogenated alkane and the above-mentioned solvents, respectively. The lotion compositions can be presented as a solution or a suspension of a compound of formula (I) in a liquid medium. A bird or mammalian host can also be protected against infestation by acarid ectoparasites by wearing a manufactured product in molded plastic of suitable form which is impregnated with a compound of formula (I). These manufactured products include collars, ear tags, bands, sheets and ribbons fixed in an adequate manner to the appropriate part of the body. Advantageously, the plastic material is a poly(vinyl chloride).

Therefore, a subject of the invention is in particular a composition containing:

- a) a compound of formula (I) as defined previously,
 - b) inert excipients suitable for use as pesticides of said product of formula (I),
- a composition containing:
- a) a compound of formula (I) as defined previously,

b) inert excipients suitable for use in the veterinary field of said product of formula (I),

and a compound of formula (I) as defined previously, for the implementation of a treatment method for the human or animal body which comprises applying a pharmaceutically acceptable formulation of said compound to said body.

The compounds of formula (I) are also useful in the protection and the treatment of plant species, in which case an effective insecticide, acaricide, molluscide or nematocide quantity of the active ingredient is applied. The application rate will vary according to the chosen compound, the nature of the composition, the method of application, the type of plant, the density of plantation, the probable infestation, and other factors, but, in general, a suitable application rate for agricultural crops is in the range of 0.001 to 3 kg per hectare, and preferably between 0.01 and 1 kg per hectare. Typical compositions for agricultural use contain between 0.0001% and 50% of a compound of formula (I) and, advantageously, between 0.1 and 15% by weight of a compound of formula (I).

The concentration of the compound of formula (I) for an application on an animal, in premises or in outside areas varies according to the chosen compound, the interval between treatments, the nature of the composition and the probable infestation, but, in general the compound must be contained in the composition applied in a quantity of 0.001 to 20.0% w/v and preferably 0.01 to 10% w/v. The quantity of compound deposited on an animal varies according to the application method, the size of the animal, the concentration of the compound in the composition applied, the dilution factor of the composition and the nature of the composition, but is generally in the range of 0.0001% to 0.5% w/w, except for undiluted compositions, such as lotion compositions which are in general deposited at a concentration in the range of 0.1 to 20.0%, and preferably 0.1 to 10%. The quantity of compound to be applied on stored products is in general in the range of 0.1 to 20 ppm. Sprays can be applied in areas so as to obtain an initial average concentration of 0.001 to 1 mg of compound of formula (I) per m³ of treated area.

12

The ointments, greases, pastes and aerosols are usually applied at random, as described above and concentrations of 0.001 to 20% w/v of a compound of formula (I) in the compound applied can be used.

The compounds of formula (I) are particularly active against lipidoptera such as *Spodoptera littoralis*, *Heliothis virescens*, *Plutella xylostella*, against coleoptera such as *Leptinotarsa decemlineata* and *Phaedon cochleariae*.

The compounds of formula (I) are thus useful for combating arthropods, for example insects and acaridae, in any environment in which they are harmful, for example in agriculture, in breeding, in public health and in domestic situations.

The harmful insects are in particular members of the orders of coleoptera (for example *Anobium*, *Ceutorrhynchus*, *Rhynchophorus*, *Cosmopolites*, *Lissorhoptrus*, *Meligethes*, *Hypothenemus*, *Hylesinus*, *Acalymma*, *Lema*, *Psylliodes*, *Leptinotarsa*, *Gonocephalum*, *Agriotes*, *Dermolepida*, *Heteronychus*, *Phaedon*, *Tribolium*, *Sitophilus*, *Diabrotica*, *Anthonomus* or *Anthrenus* spp.), lepidoptera (for example *Ephestia*, *Mamestra*, *Earias*, *Pectinophora*, *Ostrinia*, *Trichoplusia*, *Pieris*, *Laphygma*, *Agrotis*, *Amathes*, *Wiseana*, *Tryporysa*, *Diatraea*, *Sparganothis*, *Cydia*, *Archips*, *Plutella*, *Chilo*, *Heliothis*, *Spodoptera littoralis*, *Helrotuis virescens*, *Spodoptera* or *Tineola* spp.), diptera (for example *Musca*, *Aedes*, *Anopheles*, *Culex*, *Glossina*, *Simulium*, *Stomoxys*, *Haematobia*, *Tabanus*, *Hydrotaea*, *Lucilia*, *Chrysomyia*, *Callitroga*, *Dermatobia*, *Gasterophilus*, *Hypoderma*, *Hylemyia*, *Atherigona*, *Chlorops*, *Phytomyza*, *Ceratitis*, *Liriomyza* and *Melophagus* spp.), phthiraptera (*Mallophaga*, for example *Damalina* spp. and *Anoplura*, for example *Linognathus* and *Haematopinus* spp.), hemiptera (for example *Aphis*, *Bemisia*, *Phorodon*, *Aeneolamia*, *Empoasca*, *Parkinsiella*, *Pyrilla*, *Aonidiella*, *Coccus*, *Pseudococcus*, *Helopeltis*, *Lygus*, *Dysdercus*, *Oxycarenum*, *Nezara*, *Aleyrodes*, *Triatoma*, *Psylla*, *Myzus*, *Megoura*, *Phylloxera*, *Adelges*, *Nilaparvata*, *Nephrotettix* or *Cimex* spp.), orthoptera (for example *Locusta*, *Gryllus*, *Schistocerca* or *Acheta* spp.), dictyoptera (for example *Blattella*, *Periplaneta* or *Blatta* spp.), hymenoptera (for example *Athalia*, *Cephus*, *Atta*, *Solenopsis* or *Monomorium* spp.), isoptera (for example *Odontotermes* and *Reticulitermes* spp.), siphonaptera (for example *Ctenocephalides*

13

or *Pulex* spp.), thysanura (for example *Lepisma* spp.), dermaptera (for example *Forficula* spp.), psocoptera (for example *Peripsocus* spp.) and thysanoptera (for example *Thrips tabaci*).

Harmful acaridae are in particular ticks, for example the members of the following genera: *Boophilus*, *Ornithodoros*, *Rhipicephalus*, *Amblyomma*, *Hyalomma*, *Ixodes*, *Haemaphysalis*, *Dermacentor* and *Anocentor*, and acaridae and mites such as *Acarus*, *Tetranychus*, *Psoroptes*, *Notoednes*, *Sarcoptes*, *Psorergates*, *Chorioptes*, *Eutrombicula*, *Demodex*, *Panonychus*, *Bryobia*, *Eriophyes*, *Blaniulus*, *Polyphagotarsonemus*, *Scutigerella* and *Oniscus* spp.

Nematodes which attack large plants and trees in agriculture, in forestry and in horticulture, either directly, or by spreading bacterial, viral, mycoplasmal or fungal diseases of plants, are in particular root node nematodes, such as *Meloidogyne* spp. (for example *M. incognita*); cyst nematodes, such as *Globodera* spp. (for example *G. rostochiensis*); *Heterodera* spp. (for example *H. avenae*); *Radopholus* spp. (for example *R. similis*); grassland nematodes, such as *Pratylenchus* spp. (for example *P. pratensis*); *Belonolaimus* spp. (for example *B. gracilis*); *Tylenchulus* spp. (for example *T. semipenetrans*); *Rotylenchulus* spp. (for example *R. reniformis*); *Rotylenchus* spp. (for example *R. robustus*); *Helicotylenchus* spp. (for example *H. multicinctus*); *Hemicycliophora* spp. (for example *H. gracilis*); *Criconemoides* spp. (for example *C. similis*); *Trichodorus* spp. (for example *T. primitivus*); tusk nematodes, such as *Xiphinema* spp. (for example *X. diversicaudatum*), *Longidorus* spp. (for example *L. elongatus*); *Hoplolaimus* spp. (for example *H. coronatus*); *Aphelenchoides* spp. (for example *A. ritzemabosi*, *A. besseyi*); and bulb nematodes, such as *Ditylenchus* spp. (for example *D. dipsaci*).

The compounds of the invention can be combined with one or more other active pesticides constituents (for example pyrethroids, carbamates and organophosphates) and/or with attractants, repellents, bactericides, fungicides, nematocides, anthelmintics and similar products. Furthermore, it has been observed that the activity of the compounds of the invention can be improved by the addition of a synergic or potentiating agent, for example a synergic agent of the

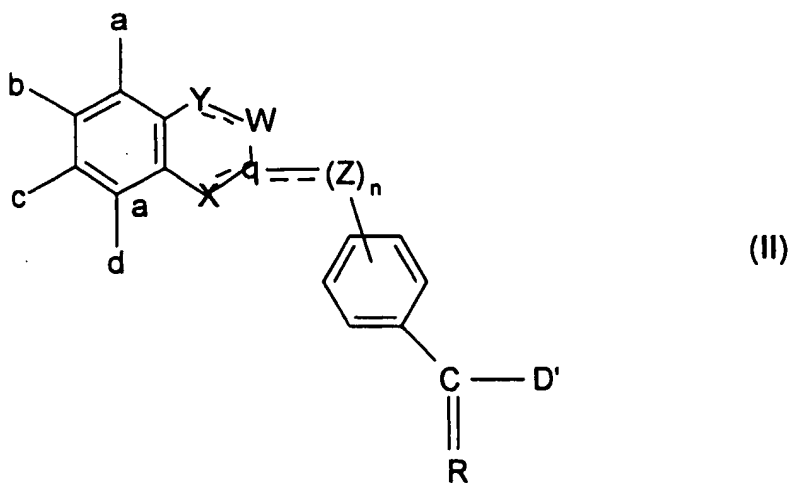
14

class of oxidase inhibitors, such as piperonyl-butoxide, or propyl 2-propynylphenyl phosphonate, by the addition of a second compound of the invention or a pyrethroid pesticide. When an oxidase-inhibiting synergic agent is present in a composition of the invention, the ratio of the synergic agent to the compound of formula (I) is in the range of 25:1 to 1:25, for example approximately 10:1.

Stabilizers used to prevent any chemical decomposition of the compounds of the invention are in particular, for example, anti-oxidants (such as tocopherols, butylhydroxy-anisole and butylhydroxytoluene), vitamin C (ascorbic acid) and oxygen trapping agents (such as epichlorhydrin) similarly organic or mineral bases, for example trialkylamines, such as triethylamine, which can act as basic stabilisers and trapping agents.

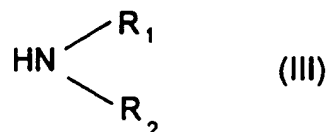
The compounds of the present invention have increased pesticide properties and photostability and/or a reduced toxicity for mammals.

A subject of the invention is also a preparation process for the compounds of formula (I), wherein a compound of formula (II):



15

in which a, b, c, d, X, Y, W, Q, Z, n and R retain their previous meaning and D' represents a hydroxy radical, a halogen atom, an alkoxy group containing up to 4 carbon atoms or a -P(O)(O ϕ) NH ϕ group in which ϕ represents a phenyl group, is subjected to the action of a compound of formula (III):



in which R₁ and R₂ retain their previous meaning in order to obtain the corresponding compound of formula (I), which is modified, if desired, in order to obtain another product of formula (I).

The products of formula (I) thus obtained can be, if appropriate, separated into their optically active isomers.

The products of formula (II) are prepared, for example, according to processes described hereafter in the experimental part, starting, e.g., from products described by M. Elliott et al., Pest. Sce 1989, 26199 (called ref. 1 in the experimental part), or by L.A. Cornelius et al., Syn. Comm. 1994, 24 (10) 2777 (called hereafter ref. 2), or starting from products prepared according to methods indicated by these authors.

Separation of the isomers can be carried out according to methods known to a person skilled in the art for example by crystallization or by chromatography.

The amidification reaction is in general carried out at a temperature comprised between -25°C and 150°C in an anhydrous and aprotic solvent such as ether, dichloromethane, toluene or benzene.

Workup and purification can be carried out by routine methods well known to s.o. skilled in the art.

16

The compounds of formula (II) used as starting material are new and are in themselves a subject of the present invention.

The disclosures in French patent application No. 97 01 541 from which this application claims priority, and in the abstract accompanying this application are incorporated herein by reference.

The following examples illustrate the invention without however limiting it.

Example 1: 4-[(6-methoxy-3,4-dihydro-2-naphthalenyl) methyl]-N-(2-methylphenyl) benzamide

0.14 ml of orthotoluidine was added at 20°C to a solution containing 0.65 ml of a 2M solution of trimethyl-aluminium in hexane and 5 ml of toluene. The solution obtained was agitated for 15 minutes at 20°C and 308 mg of the product of preparation 6 in solution in 20 ml of toluene was added. The reaction medium was taken to reflux for three hours, the temperature was then taken to 20°C and the reaction medium was treated with a solution of sodium acid phosphate. Extraction was carried out with methylene chloride followed by washing with a normal solution of hydrochloric acid, drying and concentrating. 420 mg of a product was obtained which was chromatographed on silica eluting with a heptane/dioxan mixture (7/3). In this way 351 mg of sought product was obtained.

M.p. = 162°C.

Example 2: 4-[(5,8-dibromo-2-naphthalenyl) methyl]-N-(2-methylphenyl) benzamide

Stage A: : 4-[(5,8-dibromo-2-naphthalenyl) methyl benzoic acid chloride

3 drops of DMF (N,N-dimethylformamide) were added to a solution containing 0.90 g of 4-[(5,8-dibromo-2-naphthalenyl) methyl benzoic acid in 10 ml of methylene chloride. Then 0.22 ml of oxalyl chloride was added. Agitation was carried out for 2

17

hours at ambient temperature followed by concentrating. The product obtained was taken up in 6 ml of methylene chloride. In this way a solution of the acid chloride was obtained which was used as it was in the following stage.

Stage B: 4-[(5,8-dibromo-2-naphthalenyl) methyl]-N-(2-methylphenyl) benzamide

0.20 ml of 2-methylphenylamine was introduced into 5 ml of methylene chloride. 0.26 ml of triethylamine was added. 3 ml of the solution of the acid chloride obtained in Stage A was added dropwise. The reaction medium was left under agitation for 4 hours at ambient temperature. The reaction mixture was poured into a normal solution of hydrochloric acid followed by extraction with methylene chloride, washing with a normal solution of hydrochloric acid, with a 0.5 N solution of soda, with a saturated ammonium chloride solution, filtering and concentrating. The product obtained was recrystallized from isopropyl ether followed by filtering and washing with diisopropyl ether then with pentane. The product obtained was dried under reduced pressure at 40°C. In this way 0.42 g of sought product was obtained.

Example 3: 4-[(5,6-dichloro-1-fluoro-1,2,3,4-tetrahydro-2-naphthalenyl) methyl]-N-(2-methylphenyl) benzamide

A solution containing 0.35 g of 4-[(5,6-dichloro-1-hydroxy-1,2,3,4-tetrahydro-2-naphthalenyl) methyl]-N-(2-methylphenyl) benzamide and 10 cm³ of methylene chloride is cooled down to -65°C ± 5°C. 142 mg of DAST (diethylaminosulfur trifluoride) was added. The solution was maintained under agitation for 1 hour at -65°C. The temperature was allowed to rise to 20°C and the reaction medium was poured into an aqueous solution of potassium acid carbonate, followed by agitation for 30 minutes and extraction with methylene chloride. The organic phases were collected, dried, filtered and concentrated. In this way 340 mg of sought product was obtained.

Example 4: 4-[(5,8-dibromo-2-naphthalenyl) fluoromethyl]-N-(2-methylphenyl) benzamide

0.27 g of 4-[(5,8-dibromo-2-naphthalenyl) hydroxy-methyl]-N-(2-methylphenyl) benzamide was introduced into 25 ml of methylene chloride. The reaction medium was cooled down to -15°C. 0.1 ml of DAST was added dropwise and the reaction mixture was maintained under agitation for 4 hours. A product was obtained which was poured into a saturated solution of sodium bicarbonate, agitation was carried out at ambient temperature for 45 minutes followed by extraction with methylene chloride. The organic phases were collected, followed by washing with a saturated solution of sodium chloride, drying, filtering and evaporating to dryness.

0.3 g of product was obtained which was crystallized from a pentane/ethyl acetate mixture 7-3. After filtering and washing with a pentane/ethyl acetate mixture 9-1, then with pentane, 0.21 g of sought product was obtained.

Preparation 1: methyl 4-[(3,4-dihydro-6-methoxy-1-oxo-2(1H)-naphthalenyldene) methyl] benzoate

5 g of 3,4-dihydro-6-methoxy-2(1H)-naphthalenone was dissolved in 80 ml of methanol. 4.65 g of methyl 3-formyl benzoate was added. 0.32 g of soda was added to the solution obtained, which was agitated for 66 hours at ambient temperature followed by separating, washing with methanol then with pentane and drying at 40°C for 18 hours. In this way 6.36 g of sought product was obtained. M.p. = 156°C.

Preparation 2: methyl 4-[(7-bromo-1,2,3,4-tetrahydro-naphthalenyl) methyl] benzoate

1 g of methyl 4-[(7-bromo-3,4-dihydro-1-oxo-2(1H)-naphthalenyldene) methyl] benzoate was dissolved at 20°C in 10 cm³ of chloroform. 4 cm³ of trifluoroacetic acid and 3 cm³ of triethylsilane were added. The reaction mixture was taken to reflux for 3 hours and 3 cm³ of Et₃SiH then 3 cm³ of trifluoroacetic acid were added.

19

The reaction medium was heated for 5 hours under reflux and maintained under agitation at 20°C for 48 hours. The reaction mixture was poured into a mixture of water and ice and adjusted to a basic pH by the addition of sodium acid carbonate followed by extraction with methylene chloride, drying over magnesium sulfate and filtering then concentrating under reduced pressure at 40°C. The product obtained was chromatographed eluting with a heptane/ethyl acetate mixture 9-1. 488 mg of sought product was obtained. $rf = 0.3$.

154 mg of methyl 4-[(7-bromo-3,4-dihydro-2-naphthalenyl) methyl] benzoate $rf = 0.25$, and 210 mg of methyl 4-[(7-bromo-3,4-dihydro-1-oxo-2(1H)-naphthalenyl)] benzoate $rf = 0.1$

Preparation 3: methyl 4-[(8-bromo-2-naphthalenyl) methyl] benzoate

A mixture of 537 mg of methyl 4-[(8-bromo-3,4-dihydro-2-naphthalenyl) methyl] benzoate and 48 mg of sulfur were agitated at ambient temperature while purging with nitrogen. The vessel containing the reaction mixture was immersed in a bath at 220°C for 15 minutes. The product obtained was purified by chromatography eluting with a heptane-methyl tertbutylate mixture 95-5. 266 mg of sought product was isolated.

Preparation 4: methyl 4-[(1-hydroxy-1,2,3,4-tetrahydro-6-methoxy-2-naphthalenyl) methyl] benzoate

4 g of methyl 4-[(1,2,3,4-tetrahydro-6-methoxy-1-oxo-2-naphthalenyl) methyl] benzoate and 100 ml of anhydrous methanol were mixed together and 100 ml of THF (tetrahydrofuran) was added. 0.47 g of sodium borohydride was added and the reaction medium was agitated for 2 hours 30 minutes at ambient temperature, another 0.47 g of sodium borohydride was added and agitation was continued for 30 minutes at 20°C. The reaction medium was treated with a solution of sodium acid phosphate, followed by agitation for 5 minutes, saturation with sodium chloride and extraction with ethyl acetate. After drying and concentrating the product

20

obtained was washed and dried. In this way 3.9 g of sought product was obtained melting at 112°C.

Preparation 5: methyl 4-[(1,2,3,4-tetrahydro-6-methoxy-1-oxo-2-naphthalenyl) methyl] benzoate

750 mg of 10% palladium on carbon was added to a solution containing 7.39 g of methyl 4-[(3,4-dihydro-6-methoxy-1-oxo-2(1H)-naphthalenylidene) methyl] benzoate and 200 ml of THF. The reaction medium was purged with nitrogen, with hydrogen and agitated under 1 bar of hydrogen pressure. After agitation for 2 hours the reaction medium was filtered and concentrated. 8 g of product was obtained which was purified on silica eluting with a heptane/dioxan mixture 8-2. In this way 4.92 g of sought product was isolated melting at 121°C.

Preparation 6: methyl 4-[(3,4-dihydro-6-methoxy-2-naphthalenyl) methyl] benzoate

150 mg of PTSA (p-toluene sulfonic acid) was added to a mixture containing 1.5 g of methyl 4-[(1-hydroxy-1,2,3,4-tetrahydro-6-methoxy-2-naphthalenyl) methyl] benzoate and 40 ml of toluene. The reaction medium was taken to reflux for 30 minutes, cooled down to ambient temperature, washed with a solution of sodium acid carbonate, dried and concentrated. A product was obtained which was purified by eluting with a heptane/dioxan mixture 8-2. In this way 1.35 g of sought product was isolated $rf = 0.25$.

Preparation 7: methyl 4-[(8-bromo-3,4-dihydro-2-naphthalenyl) methyl] benzoate

1.77 g of potassium *tert*-butylate was added at 20°C to a suspension containing 7.37 g of [[4-(methoxycarbonyl) phenyl] methyl] triphenyl-phosphonium bromide and 60 ml of anhydrous toluene. Agitation was carried out for 45 minutes and 2.25 g of 8-

21

bromo-3,4-dihydro-2-(1H)-naphthalenone dissolved in 12 ml of toluene was added. The reaction medium was heated to 75 ~ 80°C for 4 hours and 30 minutes, and then maintained at 20°C for 18 hours. A dilute solution of sodium acid phosphate was added and extraction was carried out with toluene. The organic phases were dried over magnesium sulfate, concentrated and 10 g of product was obtained which is chromatographed on silica eluting with a heptane/isopropyl ether mixture 8-2. In this way 1.85 g of sought product was obtained.

Preparation 8A: methyl 4-[(6-chloro-1,2,3,4-tetrahydro-2-naphthalenyl) methyl] benzoate

A mixture of 0.4 g of methyl 4-[(6-chloro-3,4-dihydro-2-naphthalenyl) methyl] benzoate, a catalytic quantity of 10% palladium on carbon and 10 ml of ethyl acetate was agitated at 25°C for 2 hours under atmospheric pressure of hydrogen, followed by separating, rinsing and bringing to dryness under reduced pressure. 0.4 g of sought product was obtained. $rf = 0.5$ heptane/ethyl acetate 7-3. $rf = 0.5$.

Preparation 8B: methyl 4-[(6-chloro-3,4-dihydro-2-naphthalenyl) methyl] benzoate

65 mg of sodium hydride at 50% in oil was added to a solution containing 240 mg of methyl 4-[(6-chloro-3,4-dihydro-2-naphthalenyl) methyl] benzoate and methyl 4-[(6-chloro-1,2,3,4-tetrahydro-2-naphthalenyldene) methyl] benzoate and 20 ml of methanol. The reaction medium was heated under reflux for 4 hours, cooled down and poured into a dilute solution of sodium acid phosphate. Extraction was carried out with methylene chloride followed by drying over magnesium sulfate and concentration. 240 mg of sought product was obtained.

22

Preparation 9: 1,1-dimethylethyl 4-[(5,8-dibromo-2-naphthalenyl) hydroxymethyl] benzoate

5,8 g of 1,1-dimethylethyl 4-iodo benzoate was introduced into 120 ml of THF. 50 mg of o-phenanthroline was added followed by cooling down to a temperature of -95/-100°C. 16 ml of a 1.6 M solution of n-butyllithium in hexane was added dropwise, then the reaction medium is maintained under agitation at -100°C for 5 minutes and a solution of 5 g of 5,8-dibromo-2-naphthalene carboxaldehyde in 20 ml of THF was added. The temperature was allowed to return to ambient and the whole was poured into a saturated solution of potassium acid phosphate, followed by extraction with ethyl ether. The ethereal phases were collected, washed with a saturated solution of sodium chloride, dried, filtered and concentrated. 11 g of a product was obtained which was purified by chromatography on silica eluting with a heptane/diisopropyl ether mixture (65-35). In this way the sought product was obtained $rf = 0.15$.

Preparation 10: 4-[(5,8-dibromo-2-naphthalenyl) methyl] benzoic acid

3.85 g of 1,1-dimethylethyl (5,8-dibromo-2-naphthalenyl) hydroxymethyl] benzoate was added to a mixture of 80 ml of acetonitrile and 80 ml of acetone. The reaction medium was cooled down to 5°C and 9.5 g of sodium iodide was added. 3.85 ml of dimethylsilyl chloride was added dropwise followed by agitation overnight at ambient temperature. The reaction medium was poured over ice, extracted with methylene chloride and ethyl ether. The organic phases were collected, washed with 1% sodium thiosulfate, with water, then with a saturated solution of sodium chloride, followed by drying, filtering and concentrating. 6.3 g of a product was obtained which was chromatographed on silica eluting with a heptane/ethyl acetate mixture 7-3 with 1% acetic acid. The sought product was obtained. $rf = 0.15$.

23

Preparation 11: 4-[(5,8-dibromo-2-naphthalenyl) acetyloxy-methyl] benzoic acid

1.5 ml of pyridine and 1 ml of acetic anhydride were added to a solution containing 1.55 g of 1,1-dimethylethyl 4-[(5,8-dibromo-2-naphthalenyl) hydroxymethyl] benzoate and 15 ml of methylene chloride. The reaction medium was maintained under agitation at ambient temperature for 2 hours. The product obtained was chromatographed on silica eluting with a heptane/diisopropyl ether mixture 6-4. The product obtained was poured into a saturated solution of potassium acid phosphate followed by extraction with methylene chloride. The organic phases were collected, washed with a 1N solution of hydrochloric acid, then with a saturated solution of sodium chloride, dried, filtered and concentrated. 2 g of a product was obtained which was chromatographed on silica eluting with a heptane/diisopropyl ether mixture 6-4. After evaporation of the fractions with an $rf = 0.25$, the product was obtained which was recrystallized from pentane. Filtering was carried out followed by washing and drying in a dessicator at 35°C under reduced pressure and 1.22 g of sought product was obtained.

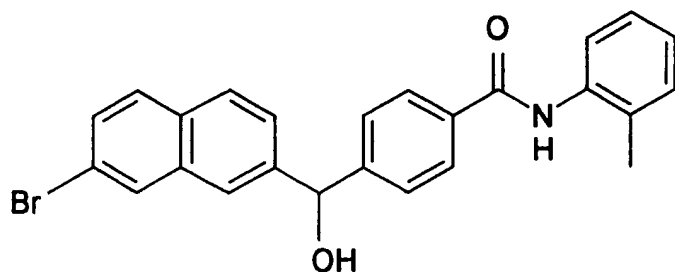
Preparation 12: 5,6-dibromo-3,4-dihydro-2(1H)-naphthalenone

4.4 ml of oxalyl chloride and a few drops of DMF were added at 0°C to a solution containing 11 g of 5,6-dibromophenylacetic acid and 100 ml of methylene chloride. The reaction medium was agitated overnight at ambient temperature and brought to dryness under reduced pressure. The product obtained was introduced into 50 ml of methylene chloride then this solution was introduced dropwise at -20°C into a mixture of 10 g of aluminum chloride and 40 ml of methylene chloride. A suspension was obtained which was kept at -20°C. A stream of ethylene was passed through it for 1 hour and 15 minutes. The reaction medium was poured over ice, extracted with methylene chloride, washed with an aqueous solution of sodium bicarbonate, dried and brought to dryness. The product obtained was chromatographed on silica eluting with a heptane/ethyl acetate mixture 70-30. The isomers obtained were

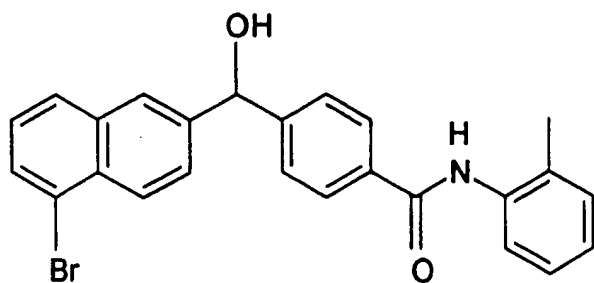
24

separated by successive crystallizations from an isopropyl ether/ methylene chloride system. In this way 30% of 5,6-dibromo-3,4-dihydro-2(1H)-naphthalenone compound was obtained, and 20% of 7,8 dibromo-3,4-dihydro-2(1H)-naphthalenone isomer was obtained.

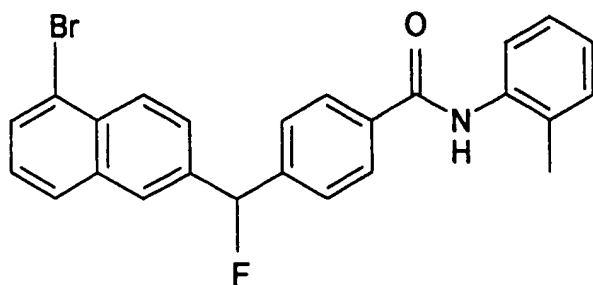
By using the processes described above the following products were prepared:



M.P. = 200°C

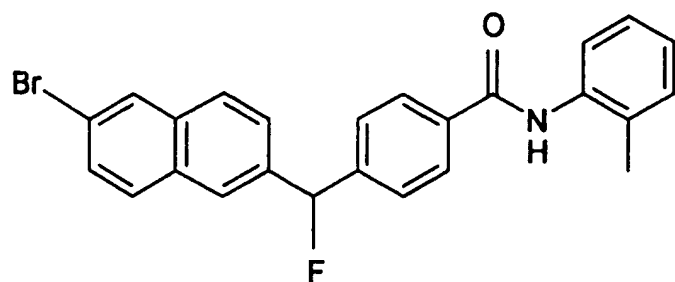


M.P. = 197°C

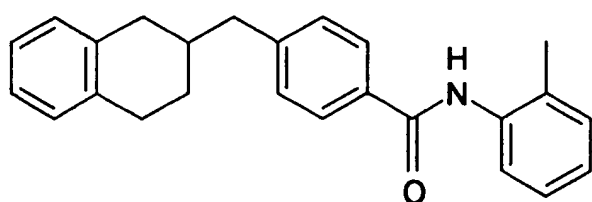


M.P. = 172°C

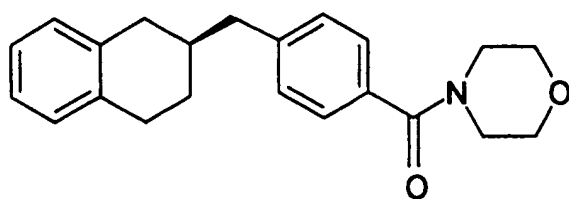
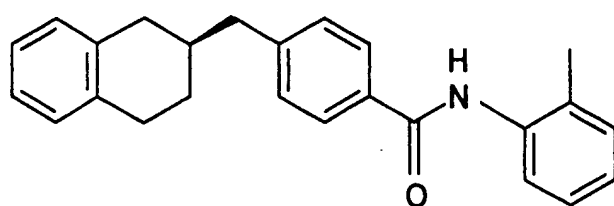
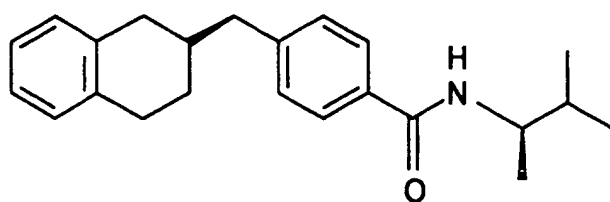
25



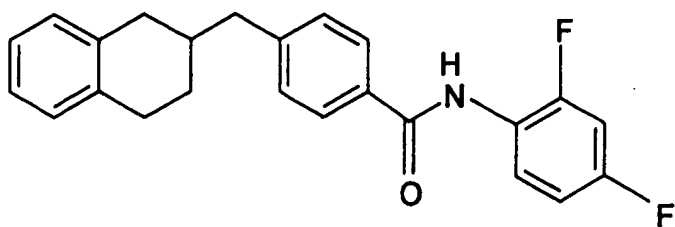
M.P. = 203°C



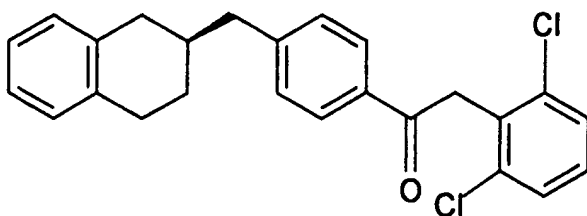
M.P. = 149°C



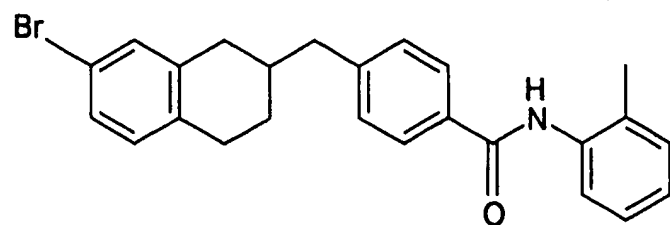
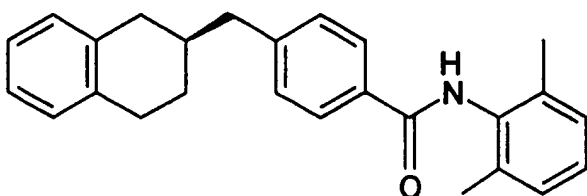
26



M.P. = 127°C

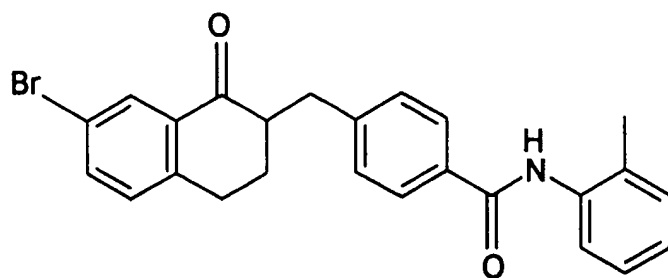


M.P. = 150°C

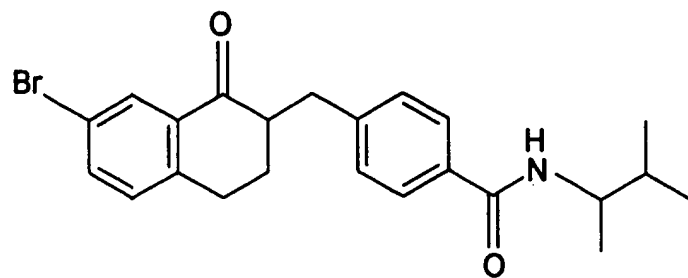


M.P. = 137°C

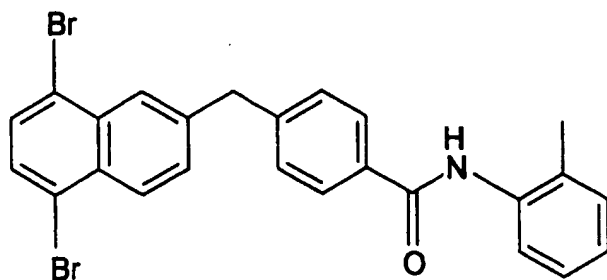
27



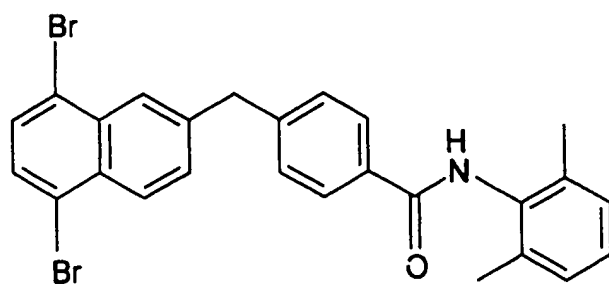
M.P. = 178°C



M.P. = 154°C

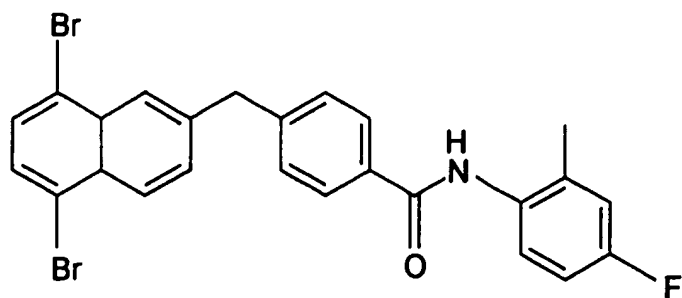


M.P. = 202°C

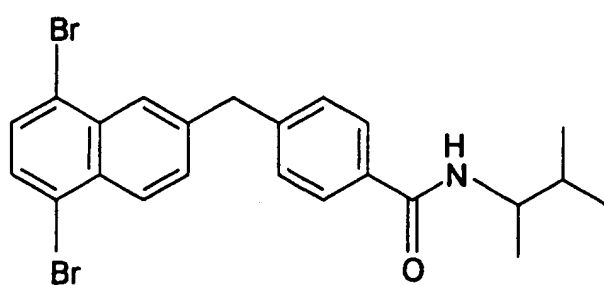


M.P. = 199°C

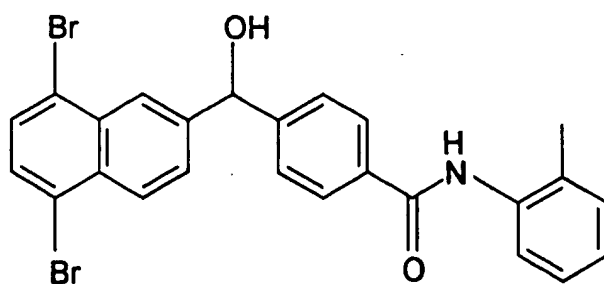
28



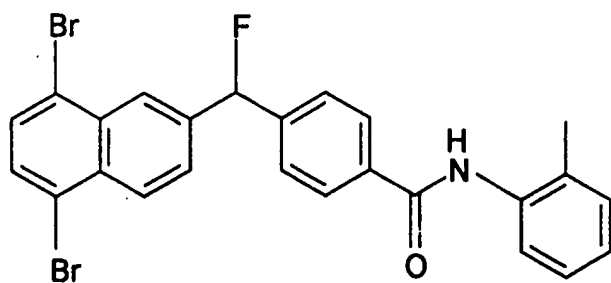
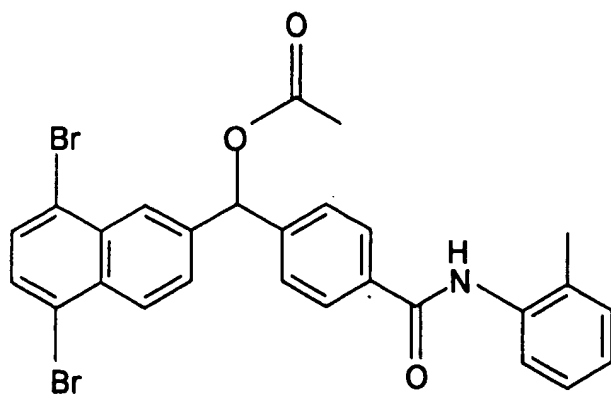
M.P. = 187°C



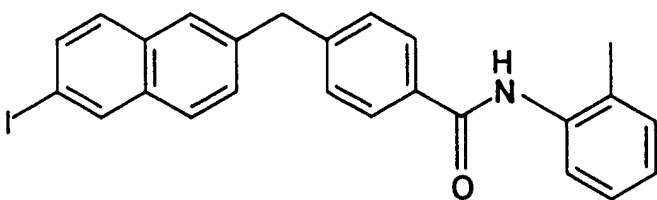
M.P. = 163°C



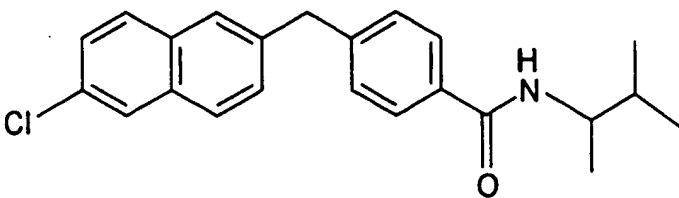
29



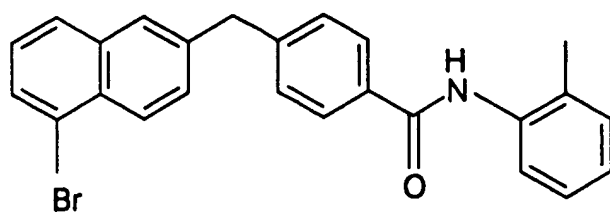
M.P. = 216°C



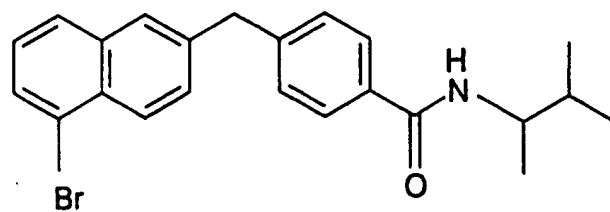
M.P. = 174°C



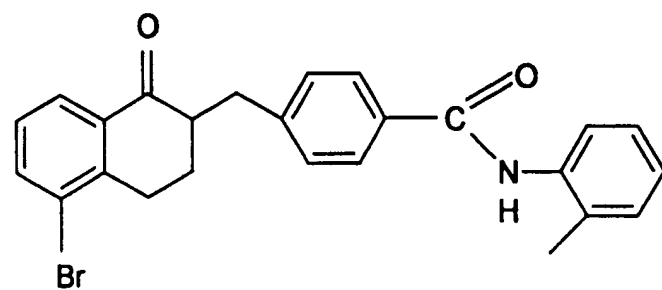
M.P. = 167°C

30

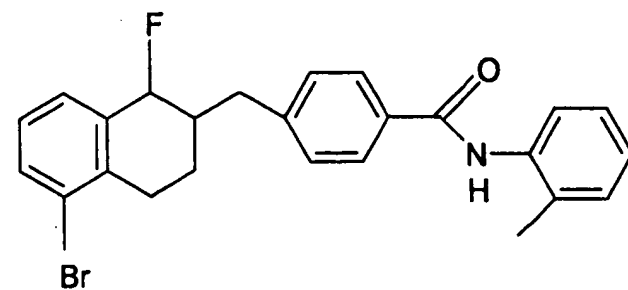
M.P. = 170°C



M.P. = 170°C

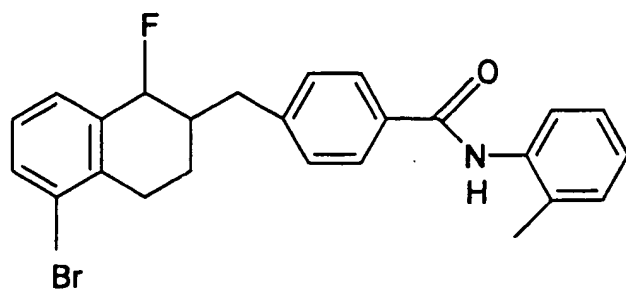


M.P. 175.2°C

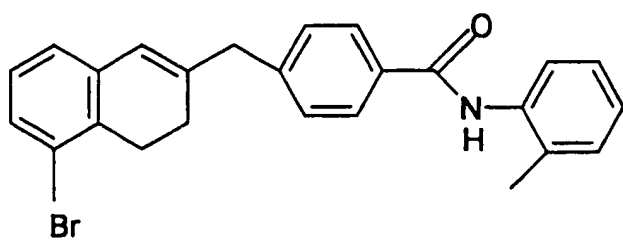


M.P. = 201.6°C

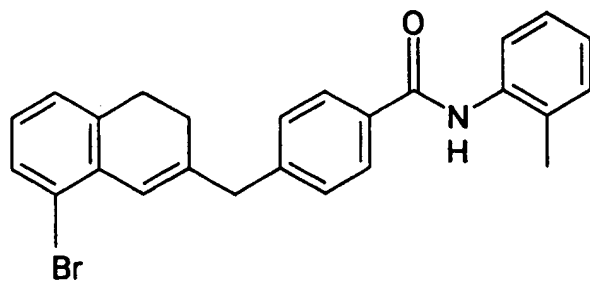
31



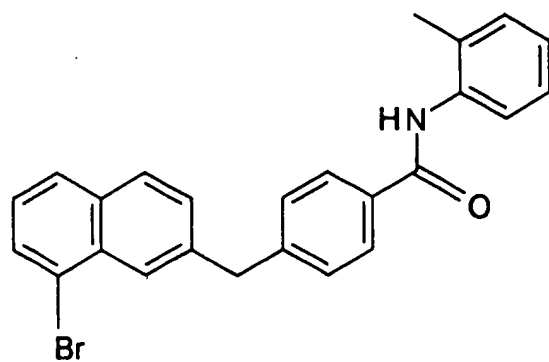
M.P. = 143.7°C



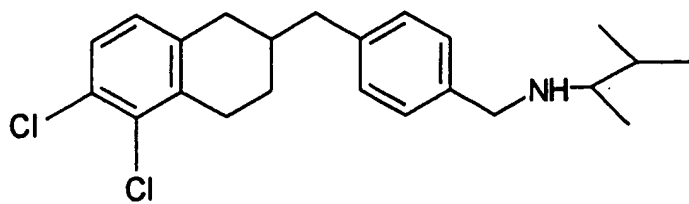
M.P. = 168.7°C



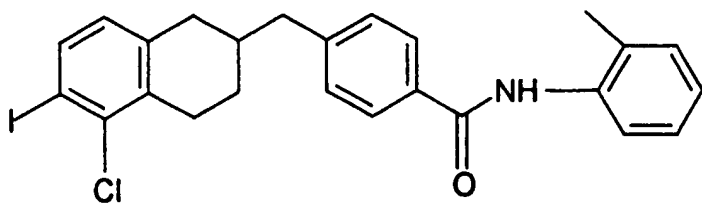
M.P. = 132°C



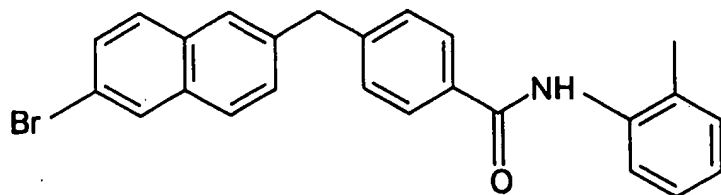
M.P. = 179°C



M.P. = 172.4°C

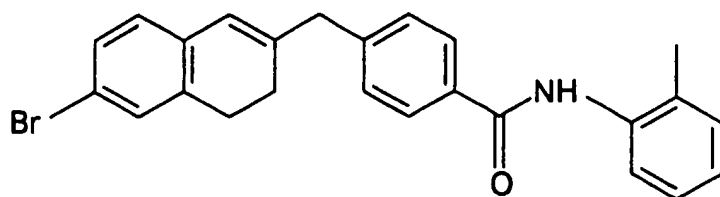


M.P. = 178.9°C

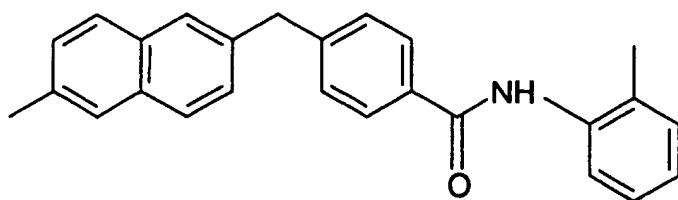


M.P. = 180.6°C

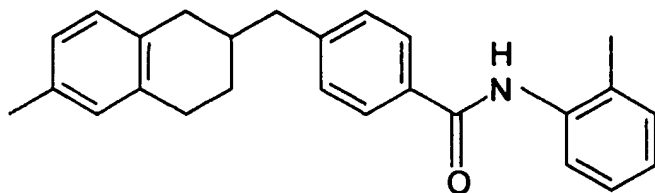
33



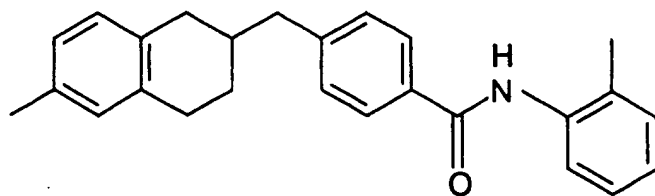
M.P. = 164.7°C



M.P. = 187.4°C

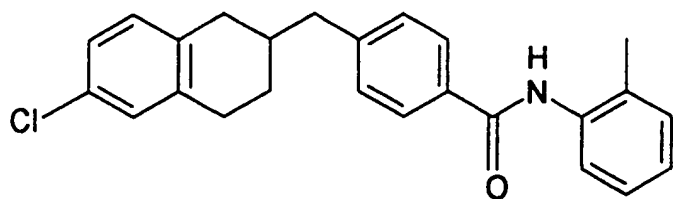


M.P. = 153.5°C

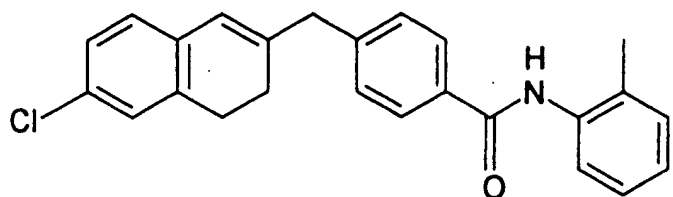


M.P. = 154°C

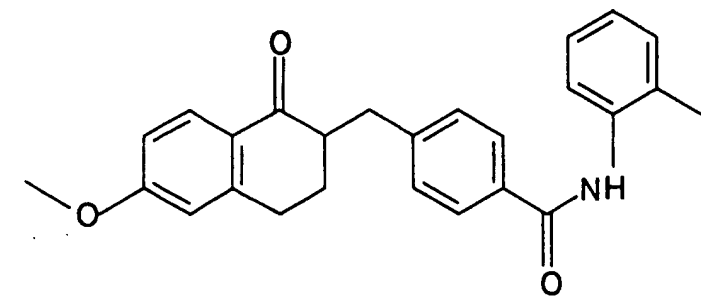
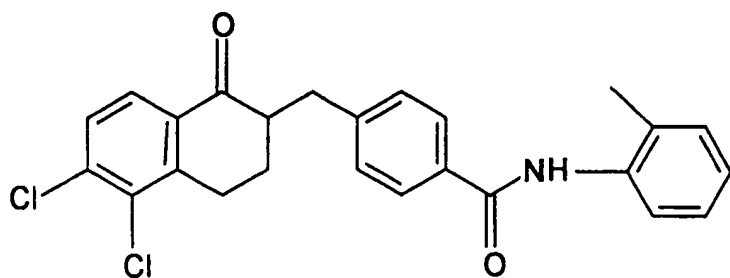
34



M.P. = 139.4°C

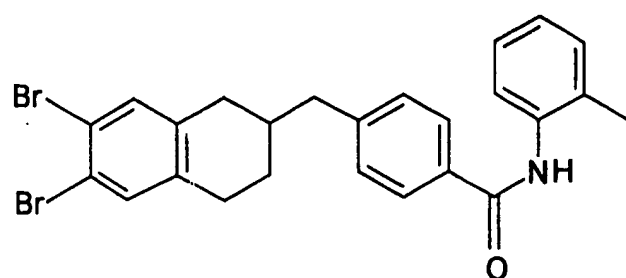
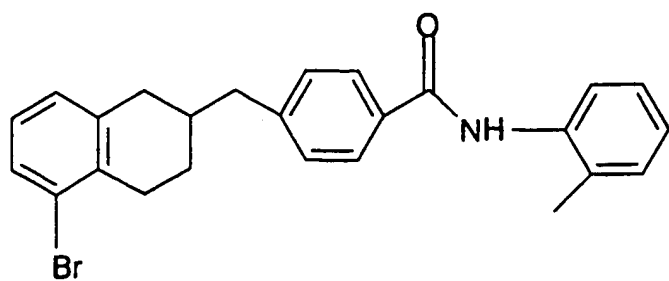


M.P. = 162°C

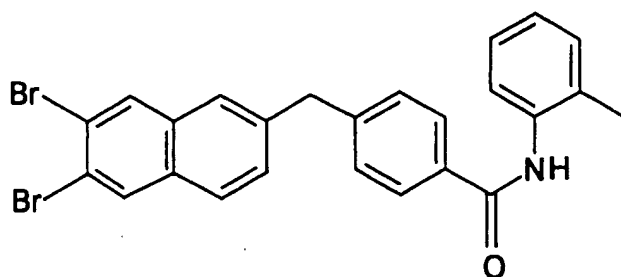


M.P. = 177°C

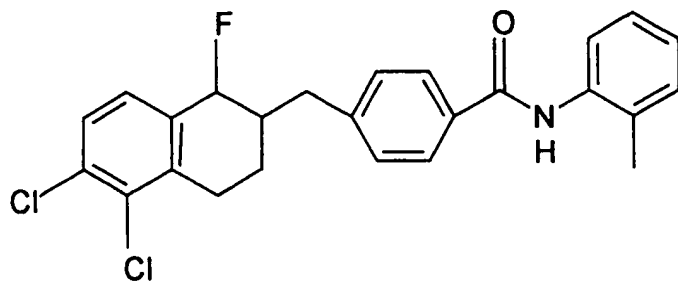
35



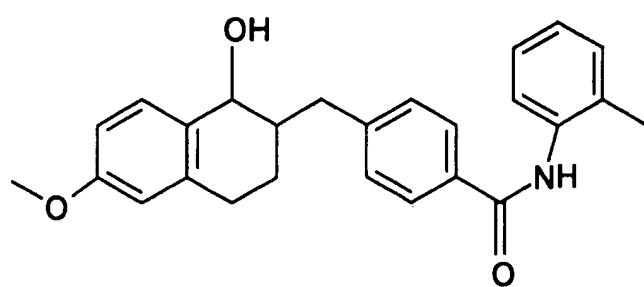
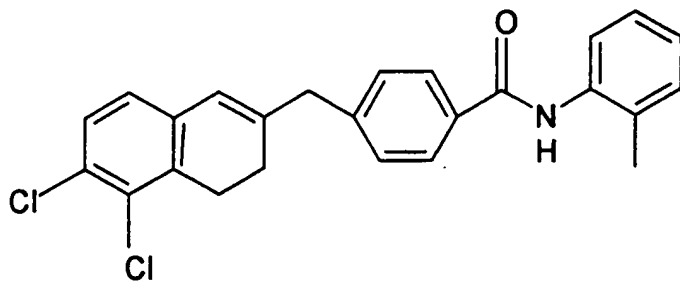
M.P. = 172°C



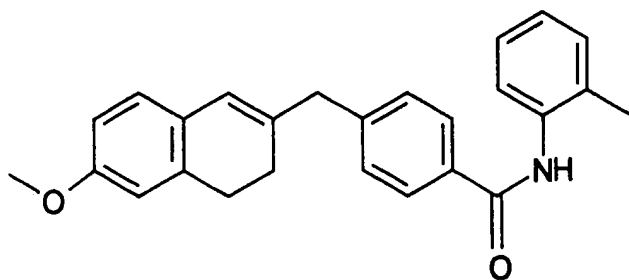
M.P. = 184°C



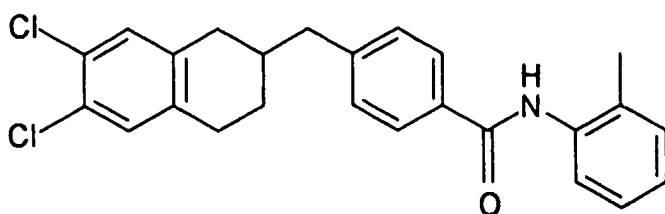
36



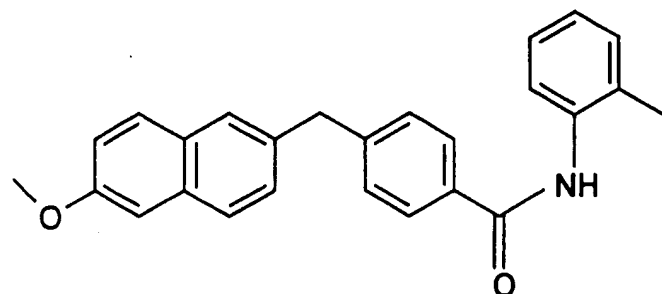
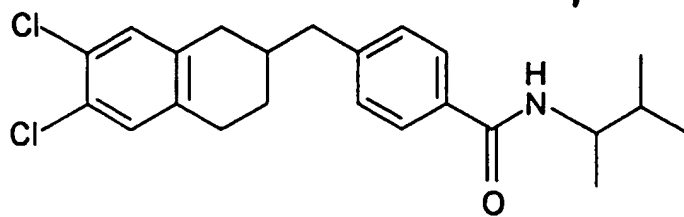
M.P. = 145°C



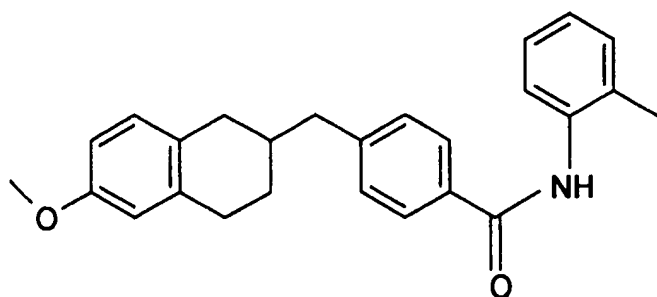
M.P. = 162°C



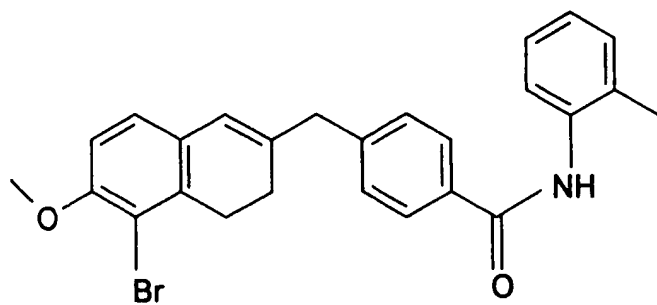
37



M.P. = 167°C

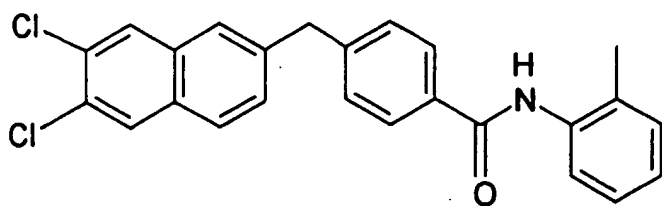


M.P. = 165°C

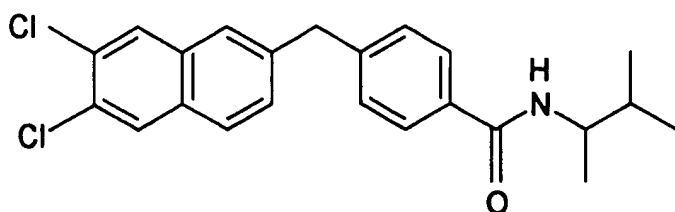


M.P. = 215°C

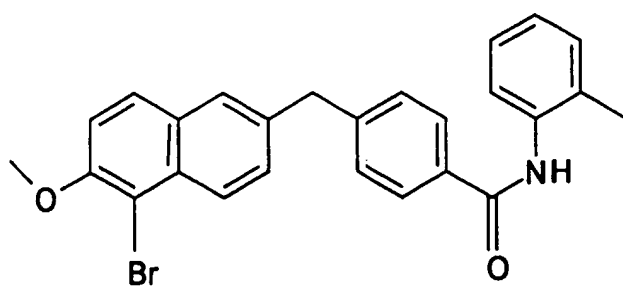
38



M.P. = 177°C



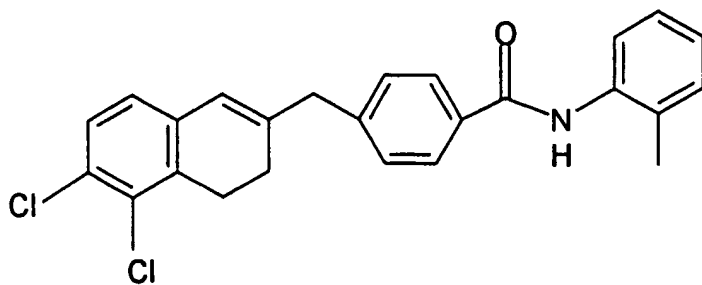
M.P. = 183°C



M.P. = 264°C

Among the products of the invention, there can in particular be mentioned the following products, which were prepared as follows:

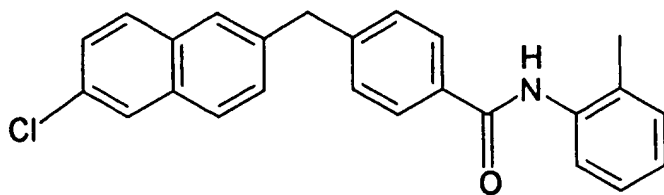
39

Product A:

rf = 0.26

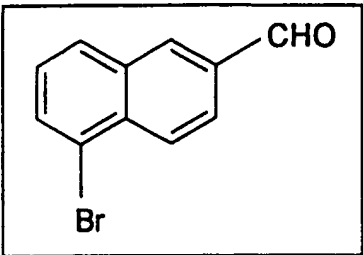
Heptane/ethyl acetate 7-3

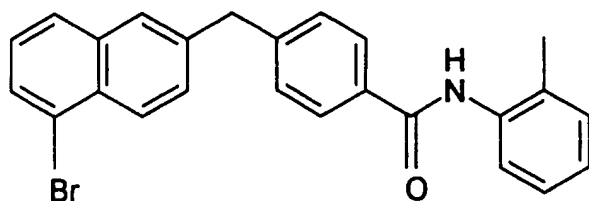
<u>Starting material:</u>	<u>Operating method:</u>
 <chem>ClC1=CC=C2C(=C(C=C1)C(=O)CCC2)Cl</chem>	Prep. 1 Prep. 2

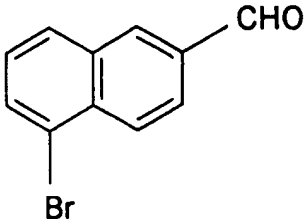
Product B:

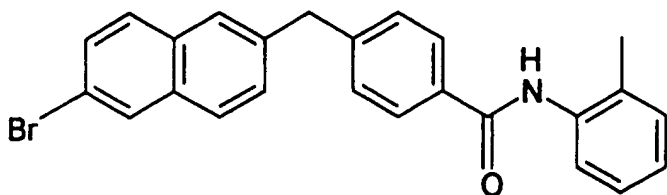
40

 $rf = 0.38$ CH_2Cl_2 -AcOEt 98/2

<u>Starting material:</u>	<u>Operating method:</u>
	Prep. Prep. 10 Ex. 2

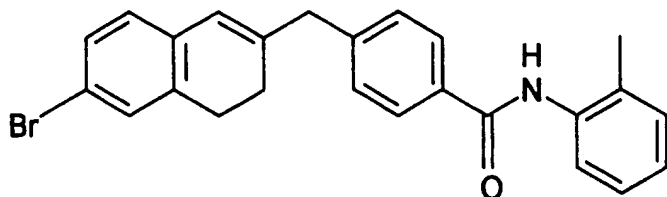
Product C: $rf = 0.33$ CH_2Cl_2 -AcOEt 98/2

<u>Starting material:</u>	<u>Operating method:</u>
	Prep. 9 Prep. 10 Ex. 2

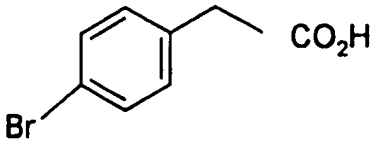
Product D:

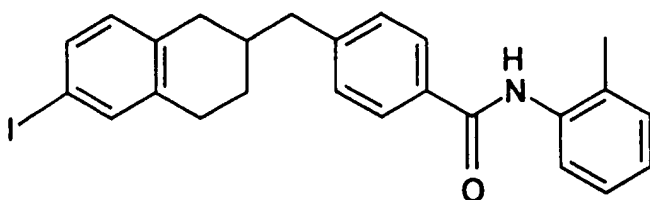
M.p. = 180.6°C

<u>Starting material:</u>	<u>Operating method:</u>
 Commercial product	Prep. 12 Prep. 7 Prep. 3 Ex. 1

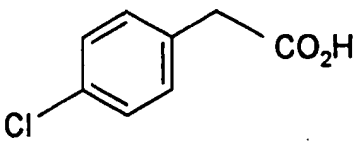
Product E:

M.p. = 164.7°C

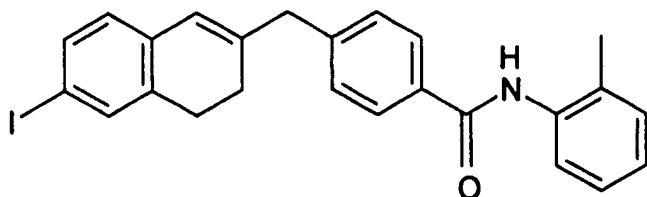
<u>Starting material:</u>	<u>Operating method:</u>
 Commercial product	Prep. 12 Prep. 7 Ex. 1

Product F:

M.p. = 139.4°C

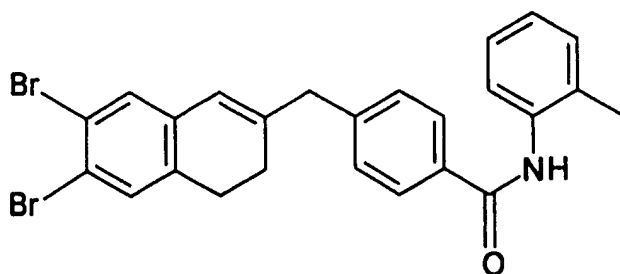
<u>Starting material:</u>	<u>Operating method:</u>
	Prep. 12 Prep. 7 Prep. 8 Ex. 1

43

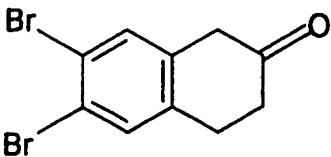
Product G:

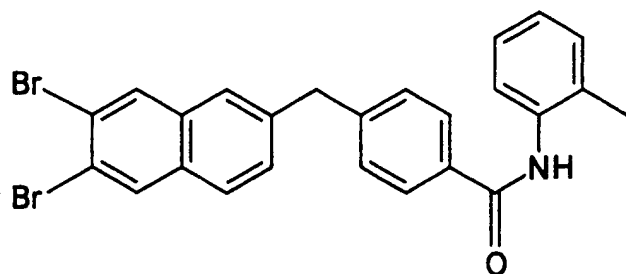
M.p. = 162°C

<u>Starting material:</u>	<u>Operating method:</u>
<p>Chemical structure of 4-chlorobenzylcarboxylic acid: A benzene ring with a chlorine atom at the 4-position and a -CH₂CO₂H group at the 1-position.</p>	Prep. 12 Prep. 7 Ex. 1

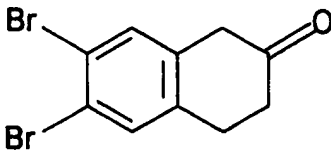
Product H:

M.p. = 172°C

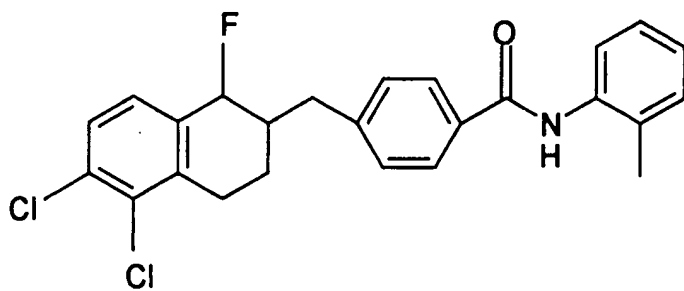
<u>Starting material:</u>	<u>Operating method:</u>
	Prep. 12 Prep. 7 Ex. 1

Product I:

M.p. = 184°C

<u>Starting material:</u>	<u>Operating method:</u>
	Prep. 12 Prep. 7 Prep. 3 Ex. 1

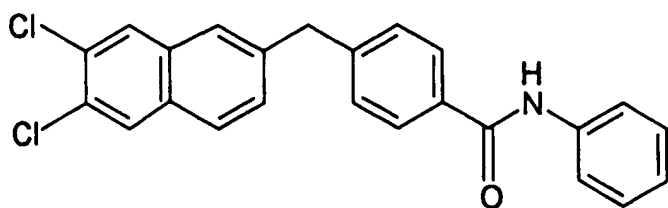
45

Product J:

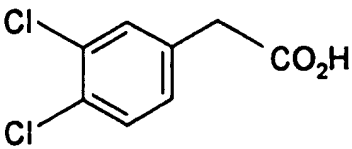
rf = 0.27

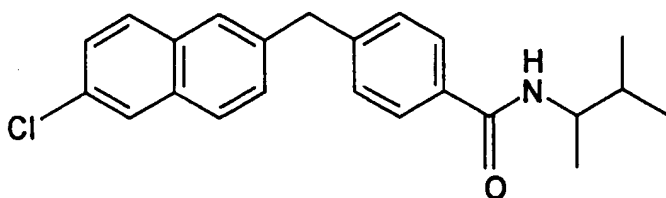
Heptane/ethyl acetate 7-3

<u>Starting material:</u>	<u>Operating method:</u>
<chem>ClC1=CC=C2C(=O)C(Cl)=CC(Cl)=CC2=C1</chem>	Prep. 1 Prep. 2 Prep. 4 Ex. 1

Product K:

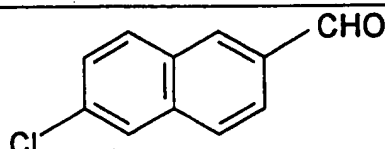
M.p. = 177°C

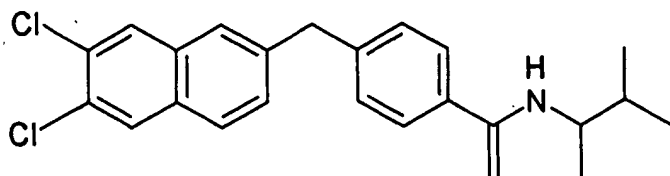
<u>Starting material:</u>	<u>Operating method:</u>
	Prep. 12 Prep. 7 Prep. 3 Ex. 1

Product L:

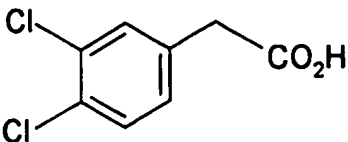
rf = 0.20

CH₂Cl₂-AcOEt 98/2

<u>Starting material:</u>	<u>Operating method:</u>
	Prep. 9 Prep. 10 Ex. 2

Product M:

M.p. = 183°C

<u>Starting material:</u>	<u>Operating method:</u>
	Prep. 12 Prep. 7 Prep. 3 Ex. 1

Preparation of compositions

In the examples of compositions below, the following signs signify:

- * Surfactant
- # Reacts by forming the polyurea walls of the microcapsule.

1. Emulsifiable concentrate.

Active ingredient	10.00
Ethoxylated alkylphenol *	7.50
Alkylarylsulfonate *	2.50
C8-C13 aromatic solvent	80.00
	—
	100.00

2. Emulsifiable concentrate.

Active ingredient	10.00
Ethoxylated alkylphenol *	2.50
Alkylarylsulfonate *	2.50
Ketonic solvent	64.00
C8-13 aromatic solvent	18.00
Antioxidant	3.00
	—
	100.00

3.	Wettable powder.	
	Active ingredient	5.00
	C8-13 aromatic solvent	7.00
	C18 aromatic solvent	28.00
	Kaolin	10.00
	Alkylarylsulfonate *	1.00
	Naphthalenesulfonic acid *	3.00
	Diatomaceous earth	46.00
		—
		100.00
4.	Dusting powder.	
	Active ingredient	0.50
	Talc	99.50
		—
		100.00
5.	Bait.	
	Active ingredient	0.5
	Sugar	79.5
	Paraffin wax	20.0
		—
		100.00
6.	Concentrate in emulsion.	
	Active ingredient	5.00
	C8-13 aromatic solvent	32.00
	Cetyl alcohol	3.00
	Polyoxyethyleneglycerol monooleate *	0.75

50

	Kaolin	25.00
		—
		100.00
10.	Granules.	
	Active ingredient	2.00
	Ethoxylated alkylphenol *	5.00
	Alkylarylsulfonate *	3.00
	C8-13 aromatic solvent	20.00
	Kieselguhr granules	70.00
		—
		100.00
11.	Aerosol (aerosol can).	
	Active ingredient	0.30
	Piperonylbutoxide	1.50
	C8-13 saturated hydrocarbonated solvent	58.20
	Butane	40.00
		—
		100.00
12.	Aerosol (aerosol can).	
	Active ingredient	0.3
	C8-13 saturated hydrocarbonated solvent	10.0
	Sorbitan monooleate *	1.0
	Water	40.0
	Butane	48.7
		—
		100.00

51

13. Aerosol (aerosol can).

Active ingredient	1.00
CO ₂	3.00
Polyoxyethyleneglycerol monooleate *	1.40
Propanone	38.00
Water	56.60
	—
	100.00

14. Lacquer.

Active ingredient	2.50
Resin	5.00
Antioxidant	0.50
Very aromatic white spirit	92.00
	—
	100.00

15. Spray (ready to use).

Active ingredient	0.10
Antioxidant	0.10
Odorless kerosene	99.80
	—
	100.00

16. Potentiated spray (ready to use).

Active ingredient	0.10
Piperonylbutoxide	0.50
Antioxidant	0.10
Odorless kerosene	99.30
	—
	100.00

52

17. Microcapsules.

Active ingredient	10.0
C8-13 aromatic solvent	10.0
Aromatic diisocyanate #	4.5
Ethoxylated alkylphenol *	6.0
Alkyldiamine #	1.0
Diethylenetriamine	1.0
Concentrated hydrochloric acid	2.2
Xanthane gum	0.2
Fumed silica	0.5
Water	64.6
	—
	100.00

18. Dispersable concentrate.

Active ingredient	5.00
N-methylpyrrolidinone	15.00
N-alkylpyrrolidinone	53.00
C8-13 aromatic solvent	16.00
Nonylphenol polyoxyethylenic ether phosphate	6.00
Ethoxylated alkylphenol	3.50
Alkylarylsulfonate	1.30
Polyalkyleneglycolic ether	0.20
	—
	100.00

19. Soluble concentrate.

A homogenous mixture is prepared of:

Active ingredient	0.25
Piperonyl butoxide	1.00

	Polyoxyethylenesorbitan esters *	0.25
	Silicon solution	0.10
	Water	58.90
		—
		100.00
7.	Concentrate in suspension.	
	Active ingredient	10.00
	Ethoxylated alkoylphenol *	3.00
	Silicon solution	0.10
	Alkanediol	5.00
	Fumed silica	0.50
	Xanthane gum	0.20
	Water	80.00
	Buffer	1.20
		—
		100.00
8.	Microemulsion.	
	Active ingredient	10.00
	Polyoxyethyleneglycerol monooleate *	10.00
	Alkanediol	4.00
	Water	76.00
		—
		100.00
9.	Granules dispersible in water.	
	Active ingredient	70.00
	Polyvinylpyrrolidone	2.50
	Ethoxylated alkylphenol	1.25
	Alkylarylsulfonate	1.25

53

Tween 80	0.25
Topanol A	0.10
Water	98.40
	<hr/>
	100.00

20. Emulsifiable concentrate.

A homogenous mixture is prepared of:

Active ingredient	0.015
Piperonyl butoxide	0.50
Topanol A	0.10
Tween 80	3.5
Xylene	95.885
	<hr/>
	100.00

STUDY OF THE BIOLOGICAL ACTIVITY**A) Study on *Phaedon cochleariae***

The product was dissolved at the desired concentration in an acetone-water mixture (50-50). Foliar disks of Chinese cabbage (*Brassica pekinensis*) were immersed for five seconds in the solution, then left to dry for one hour. Ten adults (a mixture of males and females) were added into a Petri dish containing a foliar disk. These were kept at a temperature of 25°C, with a photoperiod of twelve hours. After seven days, the mortality of the insects was checked and the foliar surface consumed is evaluated.

Product A had a very useful activity on this batch at a dose of 300 ppm.

B) Study on *Spodoptera littoralis*

The product was dissolved at the desired concentration in an acetone-water mixture (50-50). Haricot leaves (*Phaseolus vulgaris*, var. Delinel) were immersed for five seconds in the solution, then left to dry in a Petri dish for one hour. Ten larvae of *Spodoptera littoralis* were then added to each dish. These are were at a temperature of 25°C, with a photoperiod of twelve hours. After seven days, the mortality of the larvae was checked and the foliar surface consumed is evaluated.

Products A, B, C, D, E, F, G, H, I, J, K had a very good activity starting from a dose of 300 ppm.

C) Study on *Heliothis virescens*

The product was dissolved at the desired concentration in an acetone-water mixture (50-50). 50 µl of solution is deposited on the surface of small well containing approximately 2 grams of plant based artificial medium. One neonate larva of *Heliothis virescens* was then introduced into each well, which was sealed with a sheet of cellophane. The tests were kept at a temperature of 25°C, with a photoperiod of twelve hours., The mortality of the larvae was checked after seven days.

Products A, B, C, D, E, F, G, H, I, J, K, L and M had a very good activity starting from a dose of 300 ppm.

D) Study of the activity on Diabrotica

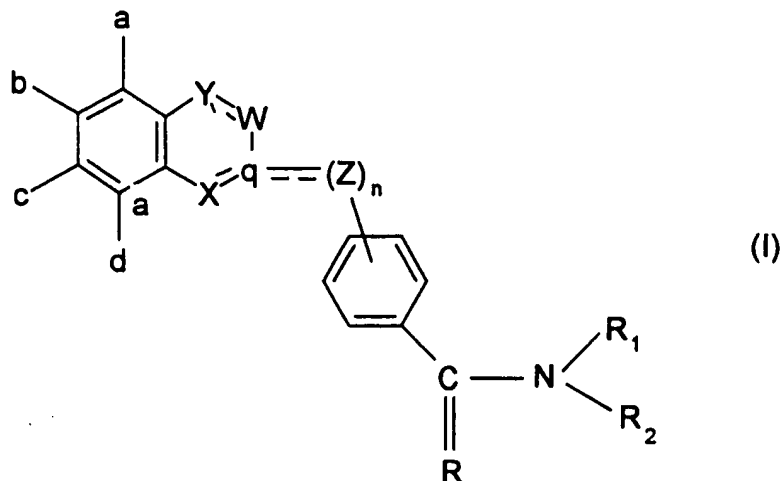
The test insects were final stage larvae of *Diabrotica undecimpunctata*.

A 9 cm disc of filter paper, placed at the bottom of Petri dish, was treated using 1 cm³ of an acetone solution. After drying 15 larvae per dose were deposited and the mortality check was carried out 24 hours after treatment.

Products A, L and M had a very good activity starting from a dose of 300 ppm.

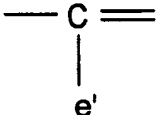
CLAIMS

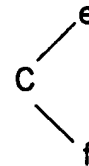
1. Compounds of formula (I):



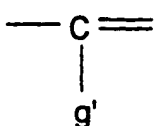
in which:

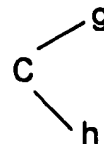
- a, b, c and d, identical to or different from one another, represent a hydrogen atom, a halogen atom, an alkyl, alkenyl or alkynyl, O-alkyl, O-alkenyl or O-alkynyl, S-alkyl, S-alkenyl or S-alkynyl radical containing up to 8 carbon atoms, optionally substituted by one or more halogen atoms, a $C\equiv N$, NO_2 or NH_2 radical, the substituents a, b, c and d being able to form between themselves rings, which either contain or do not contain one or more hetero atoms, and which are substituted or unsubstituted.
- Y and W, identical to or different from one another,

represent both  a radical or both a



radical in which e, f and e', identical or different, represent a hydrogen atom, a halogen atom, a free, etherified or esterified hydroxyl radical, or an alkyl radical containing up to 8 carbon atoms optionally substituted by one or more halogen atoms;

X represents a  a radical or a



radical in which

g, h and g', identical or different, represent a hydrogen atom, a halogen atom, a free, etherified or esterified hydroxyl radical, or an alkyl radical containing up to 8 carbon atoms optionally substituted by one or more halogen atoms, or X represents a C=O radical, an oxygen atom or a nitrogen atom, or X forms with the carbon in position 2 belonging to radical q an epoxy bridge, a cyclic hydrocarbonated radical optionally substituted by one or more halogen atoms;

- q represents a C= radical or a CD radical, in which D represents a hydrogen atom, a halogen atom or an alkyl or alkoxy radical containing up to 8 carbon atoms optionally substituted by one or more halogen atoms, or D forms with the carbon atom which carries it and one of the carbon atoms adjacent to it a carbon-carbon double bond, an epoxy radical, a cyclic hydrocarbonated radical, optionally substituted by one or more halogen atoms;

n represents an integer varying from 0 to 8,

- Z represents  a radical in which i and k, identical

or different, represent a hydrogen atom, a halogen atom, a C=N radical, a free, etherified or esterified hydroxyl radical, an SR' radical wherein R' is an organic rest containing up to 8 carbon atoms or an alkyl radical containing up to 8 carbon atoms optionally substituted by one or more halogen atoms, or Z represents an oxygen, sulfur, nitrogen atom, a C=O or C=S radical, it being understood that if n is greater than 1, Z can take different values,

- R represents an oxygen or sulfur atom;

58

- R_1 and R_2 , identical to or different from one another, represent a hydrogen atom, a linear, branched or cyclic, saturated or unsaturated, optionally substituted alkyl, CO-alkyl, CONH-alkyl or CO_2 alkyl radical, optionally interrupted by one or more preferably non-addiscent, preferably from the group consisting of N, O, S, heteroatoms, containing up to 8 carbon atoms, or an optionally substituted aryl or heteroaryl radical,

- the $-C-(Z)_n$ chain is fixed in position 3 or 4 of the benzamide, the dotted lines representing one or more optional double bonds, in all their possible isomeric forms as well as their mixtures.

2. The compounds of formula (I) defined in claim 1, in which Y represents a $-CH_2-$ radical.

3. The compounds of formula (I) defined in claim 1, in which Y and W represent a $CH=$ radical and together form a double bond in position 3(4).

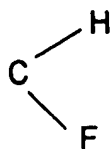
4. The compounds of formula (I) defined in any one of claims 1 or 2, in which W represents a CH_2 radical.

5. The compounds of formula (I) defined in any one of claims 1 to 4, in which q and X represent a $CH=$ radical and together form a double bond.

6. The compounds of formula (I) defined in any one of claims 1 to 4, in which q represents a $-CH-$ or $C=$ radical.

7. The compounds of formula (I) defined in any one of claims 1 to 6, in which X represents a CH_2 radical.

8. The compounds of formula (I) defined in any one of claims 1 to 6, in which X represents a



radical.

9. The compounds of formula (I) defined in any one of claims 1 to 8, in which Z represents a -CH₂- radical.
10. The compounds of formula (I) defined in any one of claims 1 to 8, in which n represents the number 1.
11. The compounds of formula (I) defined in any one of claims 1 to 10, in which R represents an oxygen atom.
12. The compounds of formula (I) defined in any one of claims 1 to 11, in which R₁ represents a hydrogen atom.
13. The compounds of formula (I) defined in any one of claims 1 to 12, in which R₂ represents an alkyl radical containing up to 8 carbon atoms or a phenyl radical optionally substituted by one or more halogen atoms and/or by one or more linear or branched alkyl radicals containing up to 8 carbon atoms.
14. The compounds of formula (I) defined in claim 13, in which R₂ represents an alkyl radical containing up to 6 carbon atoms.
15. The compounds of formula (I) defined in claim 13, in which R₂ represents a 2-methylphenyl radical.
16. The compounds of formula (I) defined in any one of claims 1 to 15, in which at least one of the substituents a, b, c and d represents a halogen atom.

17. The compounds of formula (I) defined in claim 16, in which two of the substituents a, b, c and d represent a chlorine or bromine atom.

18. The compounds of formula (I) defined in any one of claims 16 or 17, in which two of the substituents a, b, c and d represent a hydrogen atom.

19. The compounds of formula (I) defined in claim 1, the names of which follow:

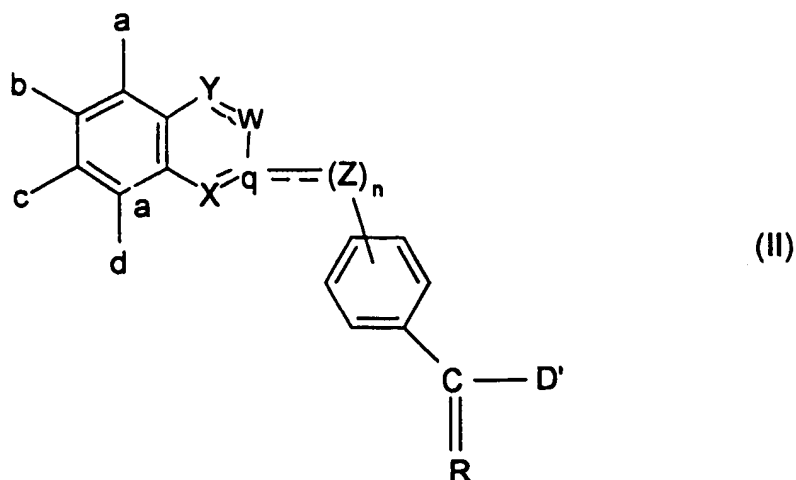
- 4-[(6-chloro-2-naphthalenyl) methyl]-N-(2-methylphenyl) benzamide,
- 4-[(5-bromo-2-naphthalenyl) methyl]-N-(2-methylphenyl) benzamide,
- 4-[(6-bromo-2-naphthalenyl) methyl]-N-(2-methylphenyl) benzamide,
- 4-[(6-bromo-3,4-dihydro-2-naphthalenyl) methyl]-N-(2-methylphenyl) benzamide,
- 4-[(6-chloro-1,2,3,4-tetrahydro-2-naphthalenyl) methyl]-N-(2-methylphenyl) benzamide,
- 4-[(6-chloro-3,4-dihydro-2-naphthalenyl) methyl]-N-(2-methylphenyl) benzamide,
- 4-[(6,7-dibromo-3,4-dihydro-2-naphthalenyl) methyl]-N-(2-methylphenyl) benzamide,
- 4-[(6,7-dibromo-2-naphthalenyl) methyl]-N-(2-methylphenyl) benzamide,
- 4-[(5,6-dichloro-1-fluoro-1,2,3,4-tetrahydro-2-naphthalenyl) methyl]-N-(2-methylphenyl) benzamide,
- 4-[(5,6-dichloro-3,4-dihydro-2-naphthalenyl) methyl]-N-(2-methylphenyl) benzamide,
- 4-[(6,7-dichloro-1,2,3,4-tetrahydro-2-naphthalenyl) methyl]-N-(2-methylphenyl) benzamide,
- 4-[(6,7-dichloro-2-naphthalenyl) methyl]-N-(2-methylphenyl) benzamide,
- 4-[(6,7-dichloro-2-naphthalenyl) methyl]-N-(1,2-dimethylpropyl) benzamide,
- 4-[(6-chloro-2-naphthalenyl) methyl]-N-(1,2-dimethylpropyl) benzamide.

20. Pesticide compositions containing at least one of the compounds defined in any one of claims 1 to 18 as active ingredient.

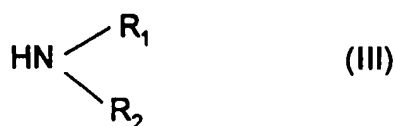
21. Pesticide compositions containing at least one compound defined in claim 19 as active ingredient.

61

22. A preparation process for the compounds of formula (I) defined in any one of claims 1 to 19 wherein a compound of formula (II):



in which a, b, c, d, X, Y, W, Q, Z, n and R are defined as in formula (I) in claim 1 and D' represents a hydroxy radical, a halogen atom, an alkoxy radical containing up to 4 carbon atoms or a -P(O)(Oφ)NHφ group in which φ represents a phenyl group is subjected to the action of a compound of formula (III):



in which R₁ and R₂ are defined as in formula (I) in claim 1 in order to obtain the corresponding compound of formula (I),

23. Compounds of formula (II) defined in claim 22.

24. The compounds of formula (II) as defined in claim 23, in which D represents an alkoxy radical containing up to 4 carbon atoms.

INTERNATIONAL SEARCH REPORT

Int. J. Application No

PCT/EP 98/02014

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C07C233/65 A01N37/22 C07C235/42 C07C235/84 C07C63/68

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C07C A01N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	DE 43 19 887 A (HOECHST SCHERING AGREVO GMBH) 22 December 1994 see claims; examples	1-24
A	EP 0 763 523 A (TEIJIN LTD) 19 March 1997 see page 25; claims	1-24



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

- "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- "&" document member of the same patent family

Date of the actual completion of the international search

7 July 1998

Date of mailing of the international search report

14/07/1998

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Sánchez García, J.M.

INTERNATIONAL SEARCH REPORT

information on patent family members

Int. l. Application No

PCT/EP 98/02014

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
DE 4319887 A	22-12-1994	AU 7123994 A	03-01-1995
		CN 1128019 A	31-07-1996
		WO 9429267 A	22-12-1994
		EP 0703899 A	03-04-1996
		HU 73352 A	29-07-1996
		JP 8511772 T	10-12-1996

EP 0763523 A	19-03-1997	AU 687202 B	19-02-1998
		AU 2538595 A	21-12-1995
		CA 2190992 A	07-12-1995
		WO 9532943 A	07-12-1995
